

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: August 30, 2005, 02:36:38 ; Search time 2069 Seconds
(without alignments)
8548.165 Million cell updates/sec

Title: US-08-731-499-9_COPY_10001_10365

Perfect score: 365

Sequence: 1 TTTGTGCTCTCCAGGCTT.....GATGCACTCCACCAAGCTTG 365

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl.*

1: gb_ba.*

2: gb_htg.*

3: gb_in.*

4: gb_om.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

8: gb_pl.*

9: gb_pr.*

10: gb_ro.*

11: gb_sts.*

12: gb_sy.*

13: gb_un.*

14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	365	100.0	10365	6	BD085733 Genes fro
2	365	100.0	121143	9	AF1312915 Homo sapi
3	365	100.0	128871	9	AL157838 Human DNA
4	363.4	99.6	105023	9	AL116668 Trypanoso
5	164	44.9	109710	9	AL050402 Human DNA
6	159.2	43.6	165365	2	AC011959 Homo sapi
7	159.2	43.6	184851	9	AC100809 Homo sapi
8	159.2	43.6	321261	9	AF131216 Homo sapi
9	153.8	42.1	188141	2	AC023120 Homo sapi
10	153.4	42.0	192499	2	AC133913 Homo sapi
11	152.6	41.8	117230	9	HSJ684FL3
12	152.2	41.7	168495	2	AC084299 Homo sapi
13	152	41.6	162216	2	AC136144 Pan trogl
14	151.4	41.5	151408	2	AC010397 Homo sapi
15	151.4	41.5	155046	2	AC146346 Pan trogl
16	151.4	41.5	182368	2	AC145822 Pan trogl
17	151.4	41.5	184010	2	AC150025 Papio anu
18	151.4	41.5	196473	2	AC146345 Pan trogl
19	151.4	41.5	253038	2	AC008930 Homo sapi

c 20	151	41.4	182509	9	AC112211	Homo sapi
c 21	150.2	41.2	157267	9	AC104938	Homo sapi
c 22	150.2	41.2	170388	2	AC027750	Homo sapi
c 23	150.2	41.2	203530	9	AC025097	Homo sapi
c 24	149.4	40.9	180303	9	AL672045	Human DNA
c 25	149.4	40.9	201012	2	AC021189	Homo sapi
c 26	149.2	40.9	167869	9	AC073342	Homo sapi
c 27	149	40.8	104295	9	AC117502	Homo sapi
c 28	149	40.8	149202	2	AC022160	Homo sapi
c 29	148.8	40.8	102562	2	AC109516	Homo sapi
c 30	148.8	40.8	119050	9	AC127024	Homo sapi
c 31	148.8	40.8	153520	9	AC130324	Homo sapi
c 32	148.8	40.8	157502	2	AC145842	Papio anu
c 33	148.8	40.8	173577	2	AC146322	Papio anu
c 34	148.8	40.8	180537	2	AC023266	Homo sapi
c 35	148.8	40.8	192030	9	AC092406	Papio anu
c 36	148.6	40.7	160099	2	AC145498	Papio anu
c 37	148.4	40.7	187738	9	AL451140	Human DNA
c 38	148.4	40.7	189456	9	AC008509	Homo sapi
c 39	148	40.5	189174	9	AC092687	Homo sapi
c 40	147.6	40.4	83661	9	AP001439	Homo sapi
c 41	147.6	40.4	100000	9	AP000143	Homo sapi
c 42	147.6	40.4	110000	2	AC143327_2	Continuation (3 of
c 43	147.6	40.4	121597	9	AP000090	Homo sapi
c 44	147.6	40.4	168778	9	HS490024	Human DNA
c 45	147.6	40.4	189256	2	AL391004	Homo sapi

ALIGNMENTS

RESULT 1
BD085733
LOCUS BD085733 10365 bp DNA linear PAT 27-AUG-2002
DEFINITION Genes from the 20q13 amplicon and their uses.
ACCESSION BD085733
VERSION JP 2001524802-A/9
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 10365)
AUTHORS Gray,J.W., Collins,C.C., Hwang,S.I., Godfrey,T., Kowbel,D. and Rommens,J.
TITLE Genes from the 20q13 amplicon and their uses
JOURNAL Patent: JP 2001524802-A 9 04-DEC-2001:
THE REGENTS OF THE UNIVERSITY OF CALIFORNIA
COMMENT OS Artificial Sequence
PN JP 2001524802-A/9
PD 04-DEC-2001
PF 15-JUL-1997 JP 1998506264
PR 15-JUL-1996 US 08/680395,16-OCT-1996 US 08/731499 PR
PI JOE W GRAY, COLIN CONRAD COLLINS, SOO IN HWANG, TONY GODFREY, PI DAVID KOWBEL,
PI JOHANNA ROMMENS
PC C12N15/11.C12Q1/68.A61K48/00
CC Description of Artificial Sequence:Genomic Sequence encoding
CC ZABCl Location/Qualifiers
FH Key 1..10365
FT source /organism='Artificial Sequence'.
FT Location/Qualifiers
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/organism='synthetic construct'
/mol_type='genomic DNA'
/db_xref='taxon:32630'

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Best Local Similarity 100.0%; Pred. No. 3.4e-90;
Matches 365; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db	10061	ATTCTTTTGATTGTTTGTAGTCTTACTTTATTTTATAGAGAAAGGCTCTTGCTCCGTCATCT	10120			/rpt_family="Alu"
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Db	10121	AGATTGGAGTGCAGCGGTGTAATCATAGCTACTGTAGTCTTGAATTCCTGAGTTCAGA	10180			/rpt_family="Alu"
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Qy	181	GATCCTTCTGCTCAGCTTCCAGGTAGCTGAGACTATATGCTGCTACCATGCACGC	240			2485..5421
Db	10181	GATCCTTCTGCTCAGCTTCCAGGTAGCTGAGACTATATGCTGCTACCATGCACGC	10240			/note="similar to Homo sapiens chromosome 5 clone CTC-269F1 deposited in GenBank Accession Number AC011341.2"
						2594..2879
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Db	10241	TGATTTTAAATTTTTTTGTAGAGTGGAGTTGCCAGGCTGGTCTTTGAACCTCCTGGCC	10300			/rpt_type=dispersed
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Qy	301	TGAGGTGATCCTCTCGTTGACCTCCCAAGTATCTTGTAGACTACAGATGCACCTCCACCAC	360			/note="similar to flow-sorted chromosome 20 HindIII fragment, SC20pF17C6 deposited in GenBank Accession Number Z94682.1"
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AL157838.24"
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Query Match      100.0%; Score 365; DB 9; Length 121143;
Best Local Similarity 100.0%; Pred. No. 3.4e-90;
Matches 365; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TTTGTGGTCTCCAAGGCTTACTTAACCTCTGTGGGTTTAACTCTTAAACCTCTGTATTTT 60
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Qy 61 ATTCTTTTGATTTGTTTGTAGCTTACTTTATTTTATAGAGAAAGGCTCTTGTCCGTCACT 120
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Qy 121 AGATTGGAGTGCAGCGGTGTAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGA 180
Db AGATTGGAGTGCAGCGGTGTAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGA 97576

Qy 181 GATCTTTCGCTCAGCTCCAGGTCCTCCAGGTCAGAGTATATATGTGCTGTACATGCACAGC 240
Db GATCTTTCGCTCAGCTCCAGGTCCTCCAGGTCAGAGTATATATGTGCTGTACATGCACAGC 97516

Qy 241 TGATTTTAAATTTTGTAGATGGAGTTGCCAGGCTGGTCTTGAACCTCTCGGCC 300
Db TGATTTTAAATTTTGTAGATGGAGTTGCCAGGCTGGTCTTGAACCTCTCGGCC 97456

Qy 301 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACCTCCACCAC 360
Db TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACCTCCACCAC 97396

Qy 361 GCTTG 365
Db GCTTG 97391

RESULT 3
AL157838/c
LOCUS
DEFINITION
Human DNA sequence from clone RP4-724E16 on chromosome
20q13.12-13.32 Contains the ZNF217 gene for zinc finger protein
217, a novel gene, a putative novel gene, ESTs, GSSs, STSs and two
CpG islands, complete sequence.
ACCESSION
AL157838
VERSION
AL157838.24 GI:9588158
KEYWORDS
HTG; CpG island; zinc finger; ZNF217.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 128871)
AUTHORS
Wilson,S.
TITLE
Direct Submission
JOURNAL
Submitted (23-FEB-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
requests: clonerequest@sanger.ac.uk
COMMENT
On Jul 31, 2000 this sequence version replaced gi:9408255.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
```

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 20, constructed by the Sanger Centre Chromosome 20 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chr20>

This sequence is the entire insert of clone RP4-724E16 The true right end of clone RPS-823G15 is at 19684 in this sequence. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. RP4-724E16 is from the library RPCI-4 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>

VECTOR: pCYPAC2.

FEATURES

source

Location/Qualifiers

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    /notes="FLAM C repeat: matches 1. .133 of consensus"
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3308..3436
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14201..14507
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14764..14868
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repeat_region
16610..16901
    /note="AluSx repeat: matches 1. .292 of consensus"
repeat_region
16989..17035
    /note="MLT1C repeat: matches 420. .463 of consensus"
repeat_region
17036..17338
    /note="AluJo repeat: matches 1. .301 of consensus"
repeat_region
17339..17730
    /note="MLT1C repeat: matches 1. .420 of consensus"
repeat_region
17958..18269
    /note="AluSx repeat: matches 1. .311 of consensus"
repeat_region
18288..18315
    /note="MER91 repeat: matches 32. .59 of consensus"
repeat_region
18383..18507

```

```
/note="MIR repeat: matches 23. .148 of consensus"
18544. .18650
/note="MIR repeat: matches 137. .261 of consensus"
18799. .18820
/note="11 copies 2 mer aa 100% conserved"
18835. .19184
/note="THB1A repeat: matches 1. .354 of consensus"
19231. .19522
/note="match: STS: Em:HSAL53YF5"
19297. .19346
/note="25 copies 2 mer ca 100% conserved"
19426. .19564
/note="MIR repeat: matches 6. .147 of consensus"
19624. .19916
/note="AluX repeat: matches 1. .295 of consensus"
20005. .20204
/note="WER20 repeat: matches 1. .218 of consensus"
20215. .20317
/note="MER5A repeat: matches 4. .108 of consensus"
20410. .20541
/note="MER5B repeat: matches 1. .120 of consensus"
20544. .20687
/note="AluSg/x repeat: matches 154. .297 of consensus"
20689. .20814
/note="FLAM A repeat: matches 14. .136 of consensus"
20816. .21218
/note="Charlie4a repeat: matches 17. .427 of consensus"
21337. .21634
/note="AluY repeat: matches 1. .297 of consensus"

Query Match      100.0%; Score 365; DB 9; Length 128871;
Best Local Similarity 100.0%; Pred. No. 3.4e-90;
Matches 365; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TTTGTGCTCTCCAAGGCTTACTTAACCTCTGTGGTTTAACTCTTAACCCCTGTGTAATTT 60
Db 64003 TTTGTGCTCTCCAAGGCTTACTTAACCTCTGTGGTTTAACTCTTAACCCCTGTGTAATTT 63944

Qy 61 ATTCCTTTTGAATTTCTTGTAGTCTTACTTATTTTATTTTATAGAGAAGGCTCTTGTCCGTCATCT 120
Db 63943 ATTCCTTTTGAATTTCTTGTAGTCTTACTTATTTTATTTTATAGAGAAGGCTCTTGTCCGTCATCT 63884

Qy 121 AGATTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAGA 180
Db 63883 AGATTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAGA 63824

Qy 181 GATCCTTCTGCTCAGCTTCCAGGTAGCTGAGACTATATGTGCTGTACATGCACAGC 240
Db 63823 GATCCTTCTGCTCAGCTTCCAGGTAGCTGAGACTATATGTGCTGTACATGCACAGC 63764

Qy 241 TGATTTTAAATTTTCTTGTAGAGATGGAGTTCGCCAGGCTGGTCTTGAACCTCTGGCC 300
Db 63763 TGATTTTAAATTTTCTTGTAGAGATGGAGTTCGCCAGGCTGGTCTTGAACCTCTGGCC 63704

Qy 301 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTTAGACTACAGATGCACCTCCACC 360
Db 63703 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTTAGACTACAGATGCACCTCCACC 63644

Qy 361 GCTTG 365
Db 63643 GCTTG 63639

RESULT 4
AC116668
LOCUS
DEFINITION AC116668 105023 bp DNA linear HTG 01-MAY-2002
IN PROGRESS ***, 3 unordered pieces.
ACCESSION AC116668
VERSION AC116668.5 GI:20376999
KEYWORDS HTG; HTGS PHASE1.
SOURCE Trypanosoma brucei
ORGANISM Trypanosoma brucei
```

```
Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.
1 (bases 1 to 105023)
El-Sayed,N.M., Ghedin,E., Song,J., Larkin,C., Wanless,D., Jones,K.,
Peterson,J., Hou,L., Zhao,H., Mason,T., Militscher,J., Pai,G., Van
Aken,S., Uterback,T., Khalak,H.G., Gerard,C., Leech,V., Ullu,E.,
Melville,S., White,O., Adams,M.D., Donelson,J.E. and Fraser,C.M.
Trypanosoma brucei GUTat10.1 RPC193-45E22 BAC genomic sequence
Unpublished
2 (bases 1 to 105023)
El-Sayed,N.M., Khalak,H. and Adams,M.D.
Direct Submission
Submitted (02-APR-2002) The Institute for Genomic Research, 9712
Medical Center Dr, Rockville, MD 20850, USA
3 (bases 1 to 105023)
El-Sayed,N.M., Khalak,H. and Adams,M.D.
Direct Submission
Submitted (01-MAY-2002) The Institute for Genomic Research, 9712
Medical Center Dr, Rockville, MD 20850, USA
On May 1, 2002 this sequence version replaced gi:20340472.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 85781: contig of 85781 bp in length
* 85782 85806: gap of unknown length
* 85807 102956: contig of 17150 bp in length
* 102957 102981: gap of unknown length
* 102982 105023: contig of 2042 bp in length.

FEATURES
source
1. .105023
/organism="Trypanosoma brucei"
/mol_type="genomic DNA"
/isolate="GUTat10.1"
/db_xref="taxon:5691"
/chromosomes="V"
/clone="RPC193-45E22"

ORIGIN

Query Match      99.6%; Score 363.4; DB 2; Length 105023;
Best Local Similarity 99.7%; Pred. No. 9.4e-90;
Matches 364; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TTTGTGCTCTCCAAGGCTTACTTAACCTCTGTGGTTTAACTCTTAACCCCTGTGTAATTT 60
Db 14284 TTTGTGCTCTCCAAGGCTTACTTAACCTCTGTGGTTTAACTCTTAACCCCTGTGTAATTT 14343

Qy 61 ATTCCTTTTGAATTTCTTGTAGTCTTACTTATTTTATAGAGAAGGCTCTTGTCCGTCATCT 120
Db 14344 ATTCCTTTTGAATTTCTTGTAGTCTTACTTATTTTATAGAGAAGGCTCTTGTCCGTCATCT 14403

Qy 121 AGATTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAGA 180
Db 14404 AGATTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAGA 14463

Qy 181 GATCCTTCTGCTCAGCTTCCAGGTAGCTGAGACTATATGTGCTGTACATGCACAGC 240
Db 14464 GATCCTTCTGCTCAGCTTCCAGGTAGCTGAGACTATATGTGCTGTACATGCACAGC 14523

Qy 241 TGATTTTAAATTTTCTTGTAGAGATGGAGTTCGCCAGGCTGGTCTTGAACCTCTGGCC 300
Db 14524 TGATTTTAAATTTTCTTGTAGAGATGGAGTTCGCCAGGCTGGTCTTGAACCTCTGGCC 14583

Qy 301 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTTAGACTACAGATGCACCTCCACC 360
Db 14584 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTTAGACTACAGATGCACCTCCACC 14643

Qy 361 GCTTG 365
Db 14643 GCTTG 36365
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```

Db      14644  GCTTG 14648

RESULT 5
HSBA46E17
LOCUS   Human DNA sequence from clone RP11-46E17 on chromosome 22, complete
DEFINITION
sequence.
ACCESSION AL050402
VERSION   AL050402.16  GI:5832404
KEYWORDS  HTG.
SOURCE    Homo sapiens (human)
ORGANISM  Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Collier,R.
Direct Submission
Submitted (01-DEC-2000) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
requests: clonerequest@sanger.ac.uk
On Sep 6, 1999 this sequence version replaced gi:5791503.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em., EMBL; Sw., SWISSPROT; Tr., TREMBL; Wp., WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep
This sequence
has been finished according to sequence map criteria as follows.
An attempt is made to resolve all sequencing problems, such as
compressions and repeats, but not necessarily within known
annotated repeat sequence elements. Where the sequence is
ambiguous, there is an annotation using the 'unsure' feature key.
This sequence was generated from part of bacterial clone contigs of
human chromosome 22, constructed by the Sanger Centre Chromosome 22
Mapping Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr22
RP11-46E17 is from the library RPCI-11.1 constructed by the group
of Pieter de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VBCOR: pBACE3.6
IMPORTANT: This sequence is not the entire insert of clone
RP11-46E17 it may be shorter because we sequence overlapping
sections only once, except for a 100 base overlap.
The true left end of clone RP1-205F14p is at 109611 in this
sequence. The true right end of clone RP1-231P7p is at 100 in this
sequence.

FEATURES             Location/Qualifiers
     source           1..109710
                     /organism="Homo sapiens"
                     /mol_type="genomic DNA"
                     /db_xref="taxon:9606"
                     /chromosome="22"
                     /clone="RP11-46E17"
                     /clone_lib="RPCI-11.1"
                     103..416
repeat_region       /note="AluSg1 repeat: matches 1..306 of consensus"
repeat_region       4948..5067
                     /note="MER45 repeat: matches 1..120 of consensus"
repeat_region       6308..6764
                     /note="L1MB5 repeat: matches 5577..6034 of consensus"
misc_feature        9512..9946
                     /note="match: GSS: Em:AQ900161"
misc_feature        9706..10111
                     /note="match: STS: Em:HSB2942C1"
repeat_region       9862..9897
                     /note="18 copies 2 mer tc 94% conserved"
repeat_region       11101..11201

```

```

/note="MER45 repeat: matches 83..178 of consensus"
11498..11578
/note="MER45 repeat: matches 1..83 of consensus"
13423..13654
/note="match: GSS: Em:AQ490130"
complement(14203..14663)
/note="match: GSS: Em:AQ613979"
14712..15061
/note="match: STS: Em:G50283
match: GSS: Em:AQ044902"
complement(17350..17747)
/note="match: GSS: Em:AQ534637"
complement(17374..17760)
/note="match: GSS: Em:AQ179064"
17772..18403
/note="match: GSS: Em:AQ508370"
17795..18238
/note="match: GSS: Em:AQ268578"
19884..19939
/note="28 copies 2 mer ag 96% conserved"
24162..24201
/note="20 copies 2 mer gt 92% conserved"
30439..30918
/note="match: GSS: Em:AQ709314"
30586..30896
/note="AluYa5 repeat: matches 1..311 of consensus"
31362..31427
/note="33 copies 2 mer gg 66% conserved"
complement(33542..33960)
/note="match: GSS: Em:AQ416617"
34761..34816
/note="L1PA5 repeat: matches 5819..5875 of consensus"
34817..35039
/note="L1PA5 repeat: matches 5861..6143 of consensus"
37097..37433
/note="match: GSS: Em:AQ136413"
41848..42021
/note="PRAM repeat: matches 1..165 of consensus"
complement(42890..43566)
/note="match: GSS: Em:AQ895335"
complement(43186..43566)
/note="match: GSS: Em:AQ166907"
43363..43781
/note="L1MD2 repeat: matches 5929..7739 of consensus"
43575..44111
/note="match: GSS: Em:AQ376177"
50205..50541
/note="MER58B repeat: matches 1..341 of consensus"
50755..50870
/note="L1MB3A repeat: matches 6050..6163 of consensus"
54192..54655
/note="232 copies 2 mer aa 56% conserved"
58652..59429
/note="match: GSS: Em:AQ739794"
61986..62254
/note="match: GSS: Em:AL219668"
complement(62014..62165)
/note="match: GSS: Em:AL253608"
62025..62196
/note="match: GSS: Em:AQ299489"
complement(62073..62436)
/note="match: GSS: Em:AL003813"
62185..62463
/note="match: GSS: Em:AZ078596"
complement(62204..62509)
/note="match: GSS: Em:AL001671"
complement(62291..62446)
/note="match: STS: Em:G08353"
complement(62314..62461)
/note="match: GSS: Em:AQ744114"
complement(join(62322..62497,79689..79824))
/note="match: STS: Em:L30312"
join(62324..62452,76137..76341)

```


Insert size: 185000; agarose-fp
Insert size: 163865; sum-of-contigs
Quality coverage: 2.8 in Q20 bases; agarose-fp
Quality coverage: 3.2 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 16 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 1204: contig of 1204 bp in length
* 1205
* 1304: gap of 100 bp
* 1305
* 3250: contig of 1946 bp in length
* 3251
* 3350: gap of 100 bp
* 3351
* 6924: contig of 3574 bp in length
* 6925
* 7024: gap of 100 bp
* 7025
* 9765: contig of 2741 bp in length
* 9766
* 9865: gap of 100 bp
* 9866
* 15163: contig of 5298 bp in length
* 15164
* 18014: contig of 2751 bp in length
* 18015
* 18016
* 18114: gap of 100 bp
* 18115
* 24325: contig of 6211 bp in length
* 24326
* 24425: gap of 100 bp
* 24426
* 31010: contig of 6585 bp in length
* 31011
* 31110: gap of 100 bp
* 31111
* 39129: contig of 8019 bp in length
* 39130
* 39229: gap of 100 bp
* 39230
* 46121: contig of 6892 bp in length
* 46122
* 46221: gap of 100 bp
* 46222
* 56362: contig of 10141 bp in length
* 56363
* 56462: gap of 100 bp
* 56463
* 67851: contig of 11389 bp in length
* 67852
* 67951: gap of 100 bp
* 67952
* 79662: contig of 11711 bp in length
* 79663
* 79762: gap of 100 bp
* 79763
* 92911: contig of 13149 bp in length
* 92912
* 93011: gap of 100 bp
* 93012
* 122796: contig of 29785 bp in length
* 122797
* 122896: gap of 100 bp
* 122897
* 122897: contig of 42469 bp in length.

FEATURES
source

1. .165365
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/clone="RP11-110L10"
/clone_lib="RPC1-11 Human Male BAC"
1. .1204
/note="assembly_fragment"
1305. .3250
/note="assembly_fragment"
3351. .6924
/note="assembly_fragment"
7025. .9765
/note="assembly_fragment"
9866. .15163
/note="assembly_fragment"
15264. .18014
/note="assembly_fragment"
clone_end:T7
vector_side:right
18115. .24325
/note="assembly_fragment"
24426. .31010
/note="assembly_fragment"
31111. .39129
/note="assembly_fragment"
39230. .46121
/note="assembly_fragment"

misc_feature 46222. .56362
/note="assembly_fragment"
misc_feature 56463. .67851
/note="assembly_fragment"
misc_feature 67952. .79662
/note="assembly_fragment"
misc_feature 79763. .92911
/note="assembly_fragment"
clone_end:SP6
vector_side:left
misc_feature 93012. .122796
/note="assembly_fragment"
misc_feature 122897. .165365
/note="assembly_fragment"

ORIGIN

Query Match 43.6%; Score 159.2; DB 2; Length 165365;
Best Local Similarity 72.4%; Pred. No. 2.8e-33;
Matches 220; Conservative 0; Mismatches 83; Indels 1; Gaps 1;

QY 58 TTTATTCCTTTGATTTGTTAGTCTTACTTTATTTTAGAAGGGTCTTGTCTCGTCA 117
Db 64164 TCTTTTTTTGTTTTTCTTTCTTTGTTGTTTAAAGACAGGGTCTTGTCTGTCA 64105

QY 118 TCTAGATTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTCTCGAGTTCA 177
Db 64104 TCCAGGCTGGATGAGTGGCAATCATAGCTCACTGCAACATGGATTCTTGAGCTCA 64045

QY 178 AGAGATCCTTTGCTCCTCAGCTTCCCAGGTAGCTGAGATATATGTGC-TGCTACCATGCA 236
Db 64044 AGCAATCCTCTGCTCCTCAGCTCCTGAGTAGCTGGAGCTAGAGCTGCACACACACACC 63985

QY 237 CAGCTGATTTTAAATTTTTTTGTAGAGATGGAGTCCCGAGGTGCTTGAACCTCT 296
Db 63984 TAGCTAAATTTTAAACATTTTTTATAGAGATGGGGTGTCCCGGCTGATCTTGAACCTCT 63925

QY 297 GGCTCAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACTCCA 356
Db 63924 GGCTCAAGTGTTCCTCTGCTCAGCTCCCAAGTGTGGGATTACAGGATGAACCA 63865

QY 357 CCAC 360
Db 63864 CCAC 63861

RESULT 7
AC100809/c
LOCUS
DEFINITION Homo sapiens chromosome 8, clone CTC-493P15, complete sequence.
AC100809
ACCESSION
VERSION AC100809.9 GI:26553405
KEYWORDS HTG.
SOURCE
ORGANISM Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 184851)
Birren,B., Nusbaum,C. and Lander,E.
Homo sapiens chromosome 8, clone CTC-493P15
Unpublished
REFERENCE
1 (bases 1 to 184851)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Boguslavsky,I., Boukhgalter,B.,
Brown,A., Camarata,J., Campiano,A., Chang,J., Chazaro,B.,
Choepel,Y., Collangelo,M., Collins,S., Collamore,A., Cook,A.,
Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,
Ferreira,P., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S.,
Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-pierre,N.,
Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliiev,I., Johnson,R.,
Jones,C., Kamat,A., Karatas,A., Kells,C., Larocque,K.,
Lamazares,R., Lander,T., Lehoczy,J., Levine,R., Liu,G.,
Maclean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C.,
McCarthy,M., McEwan,P., McKernan,K., McPheeters,R., Meldrim,J.,

Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schuback, R., Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Straus, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE

JOURNAL Submitted (22-NOV-2001) Whitehead Institute/MIT Center for Genome

REFERENCE Research, 320 Charles Street, Cambridge, MA 02141, USA

AUTHORS 3 (bases 1 to 184851)

Birren, B., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhgalter, B., Camarata, J., Chang, J., Chazaro, B., Choepel, Y., Collymore, A., Cook, A., Cooke, P., DeArrellano, K., Dewar, K., Diaz, J.S., Dodge, S., Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J., Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hafez, N., Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, G., MacLean, C., Macdonald, P., Major, J., Matthews, C., McCarthy, M., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P., Roman, J., Roy, A., Schauer, S., Schuback, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE

JOURNAL Submitted (07-NOV-2002) Whitehead Institute/MIT Center for Genome

REFERENCE Research, 320 Charles Street, Cambridge, MA 02141, USA

AUTHORS 4 (bases 1 to 184851)

Birren, B., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhgalter, B., Camarata, J., Chang, J., Chazaro, B., Choepel, Y., Collymore, A., Cook, A., Cooke, P., DeArrellano, K., Dewar, K., Diaz, J.S., Dodge, S., Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J., Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hafez, N., Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, G., MacLean, C., Macdonald, P., Major, J., Matthews, C., McCarthy, M., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P., Roman, J., Roy, A., Schauer, S., Schuback, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE

JOURNAL Submitted (12-DEC-2002) Whitehead Institute/MIT Center for Genome

REFERENCE Research, 320 Charles Street, Cambridge, MA 02141, USA

COMMENT On Dec 12, 2002 this sequence version replaced gi:24757047.

All repeats were identified using RepeatMasker:

Smit, A.F.A. & Green, P. (1996-1997)

<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WITR

Web site: <http://www-seq.wi.mit.edu>

Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L21480

Center clone name: 493_P_15

----- Location/Qualifiers

1. .184851

/organism="Homo sapiens"

FEATURES

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/mol_type="genomic DNA"
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/chromosome="8"
/map="8"
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/clone_lib="CITC Human BAC"
1. .230
/rpt_family="MERS1A-int"
232. .596
/rpt_family="MERS1A"
597. .886
/rpt_family="AluSg"
887. .1091
/rpt_family="MERS1A"
complement(1266. .1328)
/rpt_family="L2"
2361. .2598
/rpt_family="L2"
2638. .2719
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Qy 178 AGAGATCCTTTCTGCCTCAGCTTCCCAGGTAGCTAGACTATATGTGC -TGCTACCAATGCA 236
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Qy 297 GGCCTGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACCTCCA 356
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Qy 357 CCAC 360
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DEFINITION AC084299 168495 bp DNA linear HTG 22-OCT-2000
SEQUENCE, 19 unordered pieces.
ACCESSION AC084299
VERSION AC084299.1 GI:10944492
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 168495)
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
TITLE Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE Waterston,R.H.
TITLE The sequence of Homo sapiens clone
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 168495)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (21-OCT-2000) Genome Sequencing Center, Washington
University of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA

COMMENT
----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H NH0374D24
----- Summary Statistics -----
Sequencing vector: M13, 100%
Sequencing vector: plasmid; 0%
Chemistry: Dye-primer ET; 100% of reads
Chemistry: Dye-terminator Big Dye; 0% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 154635 bases at least Q40
Consensus quality: 159811 bases at least Q30
Consensus quality: 162382 bases at least Q20
Insert size: 170000; agarose-fp
Insert size: 166695; sum-of-contigs
Quality coverage: 4.03 in Q20 bases; agarose-fp
Quality coverage: 4.16 in Q20 bases; sum-of-contigs
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* NOTE: This is a 'working draft' sequence. It currently
* consists of 19 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 1283: contig of 1283 bp in length
* 1284 1383: gap of unknown length
* 1384 3418: contig of 2035 bp in length
* 3419 3518: gap of unknown length
* 3519 5659: contig of 2141 bp in length
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13355: contig of 2786 bp in length
13455: gap of unknown length
16432: contig of 2977 bp in length
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68382: contig of 10656 bp in length
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78948: contig of 10466 bp in length
79048: gap of unknown length
95469: contig of 16421 bp in length
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110400: contig of 14831 bp in length
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: August 30, 2005, 02:32:53 ; Search time 369 Seconds
(without alignments)

5855.571 Million cell updates/sec

Title: US-08-731-499-9_COPY_10001_10365

Perfect score: 365

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Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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- 5: Geneseqn2001bs:*
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- 8: Geneseqn2003as:*
- 9: Geneseqn2003bs:*
- 10: Geneseqn2003cs:*
- 11: Geneseqn2003ds:*
- 12: Geneseqn2004as:*
- 13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	365	100.0	36022	11 ACN44986	Acn44986 Human gen
3	325.8	89.3	10282	2 AAV09023	Aav09023 Homo sapi
4	147.6	40.4	310268	13 ABD32548	Abd32548 Human can
C 5	147.2	40.3	50335	9 AAD58280	Aad58280 Human tum
C 6	147.2	40.3	226475	9 AAD58279	Aad58279 Human tum
C 7	146.6	40.2	1385	4 AAK70207	Aak70207 Human imm
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9	145.8	39.9	110000	13 ABD32780_2	Continuation (3 of
10	145.2	39.8	3019	12 ADQ64936	Adq64936 Novel hum
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20	142	38.9	14327	4 AAK79116	Aak79116 Human imm

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22	142	38.9	14327	8 ADA98634	Ada98634 Human sec
23	142	38.9	14327	10 ABT16938	Abt16938 Human sec
24	142	38.9	14327	10 ABZ67608	Abz67608 Human sec
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26	140.8	38.6	84607	2 AAX90847	Aax90847 Human PAC
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28	140.4	38.5	138363	13 ABD32624	Abd32624 Human can
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C 34	140	38.4	26764	2 AAT71696	Aat71696 Human deo
35	140	38.4	26865	12 ADM97421	Adm97421 Prostate
C 36	140	38.4	34739	11 ACN45078	Acn45078 Human gen
37	140	38.4	171158	12 ADQ97894	Adq97894 Human can
C 38	139.8	38.3	5230	5 AAF73808	Aaf73808 Partial h
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40	139.6	38.2	26928	5 ABA82620	Abas82620 Human HBM
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42	139.6	38.2	26928	8 ACC45361	Acc45361 Human HBM
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44	139.6	38.2	26928	10 ADB82430	Ade82430 Human DNA
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ALIGNMENTS

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DT 07-NOV-2001 (first entry)
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Human immune/haematopoietic antigen genomic sequence SEQ ID NO:35905.
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DE Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
KW Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
KW cytosstatic; gene therapy; vaccine; metastasis; ds.
XX
XX Homo sapiens.
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XX WO200157182-A2.
XX
XX 09-AUG-2001.
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XX 17-JAN-2001; 2001WO-US0001354.
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XX 31-JAN-2000; 2000US-0179065P.
PR 04-FEB-2000; 2000US-0180828P.
PR 24-FEB-2000; 2000US-0184664P.
PR 02-MAR-2000; 2000US-0186350P.
PR 16-MAR-2000; 2000US-0189874P.
PR 17-MAR-2000; 2000US-0190076P.
PR 18-APR-2000; 2000US-0198123P.
PR 19-MAY-2000; 2000US-0205515P.
PR 07-JUN-2000; 2000US-0209467P.
PR 28-JUN-2000; 2000US-0214886P.
PR 30-JUN-2000; 2000US-0215135P.
PR 07-JUL-2000; 2000US-0216647P.
PR 07-JUL-2000; 2000US-0216880P.
PR 11-JUL-2000; 2000US-0217487P.
PR 11-JUL-2000; 2000US-0217496P.
PR 14-JUL-2000; 2000US-0218290P.
PR 26-JUL-2000; 2000US-0220963P.
PR 26-JUL-2000; 2000US-0220964P.
PR 14-AUG-2000; 2000US-0224518P.
PR 14-AUG-2000; 2000US-0224519P.
PR 14-AUG-2000; 2000US-0225213P.
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Qy 121 AGATTGAGTGCACGGGTGAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGA 180
Db 6524 AGATTGAGTGCACGGGTGAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGA 6583
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Qy 301 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTTAGACTACAGATGCACCTCCACCAC 360
Db 6704 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTTAGACTACAGATGCACCTCCACCAC 6763
Qy 361 GCTTG 365
Db 6764 GCTTG 6768

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XX AC ACN44986;
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XX 18-NOV-2004 (first entry)
XX Human genomic sequence hCG37127.
XX Cytostatic; carcinoma; lymphoma; cancer; human; gene; ss.
XX Homo sapiens.
XX WO2003073826-A2.
XX 12-SEP-2003.
XX 28-FEB-2003; 2003WO-US006235.
XX 01-MAR-2002; 2002US-00087192.
XX (SAGR-) SAGRES DISCOVERY.
XX Morris DW;
XX WPI; 2003-328604/31.
XX Recombinant nucleic acid useful for diagnosis and treatment of carcinoma
XX comprises a nucleotide sequence.
XX Claim 1; SEQ ID NO 1708; Opp; English.
XX The present invention relates to novel DNA and protein sequences which
XX are associated with carcinomas. The sequences are useful for: (i) for
XX screening drug candidates; (ii) for screening of bioactive agent capable
XX of binding to Carcinoma Associated Protein (CAP); (iii) for screening of
XX a bioactive agent capable of modulating the activity of CAP; (iv) for
XX evaluating the effect of a candidate carcinoma drug; (v) for diagnosing
XX carcinoma; (vi) for inhibiting the activity of CAP; (vii) for treating

CC carcinoma; (viii) for neutralizing the effect of CAP; (ix) as a biochip;
CC (x) for diagnosing carcinoma or a propensity to carcinoma; and (xi) for
CC determining Carcinoma Associated (CA) gene copy number. In addition, the
CC CA genes are useful as DNA vaccines and the CAP are useful as markers of
CC carcinoma including lymphoma. The present sequence is one such CA coding
CC sequence. Note: This patent is an equivalent to basic patent
CC US2002182586A1, for which no sequence data was published
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Db 17891 GCTTG 17895
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XX 20q13 amplicon; chromosome 20; tumour; detection; ZABC-1 gene;
XX chromosomal abnormalities; probe; gene therapy; antisense inhibition;
XX treatment; age-related macular degeneration; retinitis pigmentation;
XX Leber's congenital amaurosis; zinc finger amplified in breast cancer; ds.
XX Homo sapiens.
XX WO9802539-A1.
XX 22-JAN-1998.
XX 15-JUL-1997; 97WO-US012343.
XX 15-JUL-1996; 96US-00680395.
XX 16-OCT-1996; 96US-00731499.
XX 17-JAN-1997; 97US-00785532.
XX (REGC) UNIV CALIFORNIA.
XX Gray JW, Collins CC, Hwang S, Godfrey T, Kowbel D, Rommens J;
PI

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 Db 77890 TTTTCTTTTCTTTTCTTTTGTAGACAGAGTCCCACTCCATCAACAGGCTGGAGTGCAGTGG 77949
 QY 138 TGTAAATCATAGCTTACCTGAGTCTTGAATTCCTGAGTTCAAGAGATCCTTCTGCCTCAGC 197
 Db 77950 TGCAGTCTTGCTCCTCAGTGCATCTTTGCACTCTGGGCTCAAGTGAATCTCGTGCCTCAGC 78009
 QY 198 TTCCAGGATGCTGAGACTATATGTGTCTGTCTACCATGCACAGCTGATTTTAAATTTTTT 257
 Db 78010 CTCCAGTAGCTGAGATTACAGGTGCCACACCACTTCTCAGATAATTTTGTATTTTTA 78069
 QY 258 TTGTAGAGATGG-----AGTTGCCAGGCTGGTCTTGAATCTCTGGCTGAGGTGAT 309
 Db 78070 CTAGAGATGGGGTTTACCATGTTGGCCAGGCTGGTCTCGAACTCTCTGACCTCAAGTGA 78129
 QY 310 CCTCTCGGTGACCTCCCAAGTATCTTAGACTACAGATGCATCCACACCGCTTG 365
 Db 78130 CCACCTGTCTTGGCTCTCCAAAGTGTGGGATTACAGGTGCGAACCACTGTGCTG 78185

RESULT 5

AAD58280/c

ID AAD58280 standard; DNA; 50335 BP.

XX AAD58280;

XX 20-NOV-2003 (first entry)

XX Human tumour suppressor gene, Lmt intron 1 DNA.

XX Tumour suppressor gene; Lmt; cancer; therapy; cytostatic; human; ds.

XX Homo sapiens.

XX PN W02003066869-A1.

XX PD 14-AUG-2003.

XX PF 07-FEB-2003; 2003WO-AU000126.

XX PR 07-FEB-2002; 2002AU-00000371.

XX PA (HALL-) HALL INST MEDICAL RES WALTER & ELIZA.

XX PI Cook WD, Mccaw BJ;

XX WPI; 2003-646311/61.

XX New nucleic acid molecule, useful for screening a subject for the presence of an aberration in a gene encoding an LMT.

XX Claim 10; Page 299-314; 373pp; English.

XX The invention relates to novel tumour suppressor gene, referred to as Lmt. The invention also relates to the field of cancer therapy and cancer diagnostics. The nucleic acid molecule is useful for screening a subject for the presence of an aberration in a gene encoding an LMT. The present sequence is human Lmt intron 1 DNA

SQ Sequence 50335 BP; 15699 A; 9846 C; 9072 G; 14424 T; 0 U; 1294 Other;

Query Match

Best Local Similarity 40.3%; Score 147.2; DB 9; Length 50335;

Matches 213; Conservative 0; Mismatches 68; Indels 5; Gaps 2;

QY 75 TTATGCTTACTTTTATTTTGTAGAGAAAGGGTCTTGTCCGTCTCATCTAGATTGGAGTGCAG 134
 Db 35888 TCTTTTTTTTTTTTTTTTTTGTAGACAAGGTCTCACTCTGTCTATCCAGGCTGAAGTGCAG 35829

QY 135 CGGTGTAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGAGATCCTTCTGCCTC 194
 Db 35828 TGGCATGATCATGGCTCAGTGTAGCTCGACCTCTCGAGCTCAGGTGATCCTCCACCTC 35769

QY 195 AGCTTCCAGGTAGCTGAGACTATATGTG-CTGCTACCATGCACAGCTGATTTTAAATT 253
 Db 35768 AGCTTCCAGGTAGCTGGGACTATAGGAGTGTGCTACCATGCCTAGCTAATTTTAAATT 35709
 QY 254 TTTTGTGTAGAGATGAGATTGCCAGGCTGGTCTTGAACCTCTGGCTGAGGTGATCCTC 313
 Db 35708 CTTTGTGTTT---GTTATGTTGCCAGGCTGGTCTCGAACTCTCTGGGCTCAGGTGATCCAC 35653
 QY 314 CTGCGTTGACCTCCCAAGTATCTTAGACTACAGATGCACATCCCAACCA 359
 Db 35652 CCGCCTTGGCTCTCCACAGCTGTGAGATTACAGACGCTGAGGCCACCA 35607

RESULT 6

AAD58279/c

ID AAD58279 standard; DNA; 226475 BP.

XX AAD58279;

XX 20-NOV-2003 (first entry)

XX Human tumour suppressor gene, Lmt reverse complement DNA.

XX Tumour suppressor gene; Lmt; cancer; therapy; cytostatic; human; ds.

XX Homo sapiens.

XX PN W02003066869-A1.

XX PD 14-AUG-2003.

XX PF 07-FEB-2003; 2003WO-AU000126.

XX PR 07-FEB-2002; 2002AU-00000371.

XX PA (HALL-) HALL INST MEDICAL RES WALTER & ELIZA.

XX PI Cook WD, Mccaw BJ;

XX WPI; 2003-646311/61.

XX New nucleic acid molecule, useful for screening a subject for the presence of an aberration in a gene encoding an LMT.

XX Claim 10; Page 233-299; 373pp; English.

XX The invention relates to novel tumour suppressor gene, referred to as Lmt. The invention also relates to the field of cancer therapy and cancer diagnostics. The nucleic acid molecule is useful for screening a subject for the presence of an aberration in a gene encoding an LMT. The present sequence is human Lmt reverse complement DNA

SQ Sequence 226475 BP; 61024 A; 41761 C; 40916 G; 57494 T; 0 U; 25280 Other;

Query Match

Best Local Similarity 40.3%; Score 147.2; DB 9; Length 226475;

Matches 213; Conservative 0; Mismatches 68; Indels 5; Gaps 2;

QY 75 TTATGCTTACTTTTATTTTGTAGAGAAAGGGTCTTGTCCGTCTCATCTAGATTGGAGTGCAG 134
 Db 211265 TCTTTTTTTTTTTTTTTTTTGTAGACAAGGTCTCACTCTGTCTATCCAGGCTGAAGTGCAG 211206

QY 135 CGGTGTAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGAGATCCTTCTGCCTC 194
 Db 211205 TGGCATGATCATGGCTCAGTGTAGCCTCGACCTCTCGAGCTCAGGTGATCCTCCACCTC 211146

QY 195 AGCTTCCAGGTAGCTGAGACTATATGTG-CTGCTACCATGCACAGCTGATTTTAAATT 253
 Db 211145 AGCTTCCAGGTAGCTGGGACTATAGGAGTGTGCTACCATGCCTAGCTAATTTTAAATT 211086

QY 254 TTTTGTGTAGAGATGAGATTGCCAGGCTGGTCTTGAACCTCTCGGCTGAGGTGATCCTC 313
 Db 211145 AGCTTCCAGGTAGCTGGGACTATAGGAGTGTGCTACCATGCCTAGCTAATTTTAAATT 211086


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Qy 352 CTCACACACG 362
Db 779 AGCCACCATGC 769

RESULT 9
ABD32780_2
Continuation (3 of 5) of ABD32780 from base 200001 (Human cancer-associated genomic DNA
WP Sequence split into 5 fragments LOCUS ABD32780 Accession Abd32780
WP Fragment Name Begin End
WP ABD32780_0 1 110000
WP ABD32780_1 100001 210000
WP ABD32780_2 200001 310000
WP ABD32780_3 300001 410000
WP ABD32780_4 400001 430442

Query Match 39.9%; Score 145.8; DB 13; Length 110000;
Best Local Similarity 67.3%; Pred. No. 9.2e-29;
Matches 239; Conservative 0; Mismatches 107; Indels 9; Gaps 2;

Qy 19 TTAATTAACTCTGGGTTTAACTCTTAACCTGTGTATTTTATCTTTTGATTGTTA 78
Db 21592 TCACCTATTCTTTTGATGTATTTATTTATTTATTTATTTATTTATTTATTT 21651

Qy 79 GTCTACTTTATTTTAGAGAAAGGGTCTTGCTCCGTCATCTAGATTGGAGTCAGCGGT 138
Db 21652 ATTTTATTTATTTTGGAGACAGAGTCTGGCTGTGTCAACCAGGCTGGAGTCAGTGGC 21711

Qy 139 GTAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGAGATCCTTCTGCTCAGCT 198
Db 21712 ACGATCTCGGCTCGCTCAACCTCTGCCTCTCTGGGTTCAAGTGAATCTCTGCTCAGCC 21771

Qy 199 TCCAGGTAGCTGAGACTATATGTGCTGTACCATGCACAGCTGATTTTAAATTTTTT 258
Db 21772 TCCAGGTAGCTGGATTACAAGCACT-CCACACAGCCGAGCTAAATTTTGTATTTTAG 21830

Qy 259 TGTAGAGATGG-----AGTTGCCAGGCTGTCTTTGAACTCTGCGCTGAGGTGATC 310
Db 21831 TGGAGATGGGTTTCACTATGTTGGCCAGGCTGTGTTGGAACTCTCTGGCTCAGTGATC 21890

Qy 311 CTCTGGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACATCCACACGCTTG 365
Db 21891 CACCTGCTTTGGCTCCCAAAGTCTGGGATTACAGGTGTGAGCCACCGCACCTG 21945

RESULT 10
ADQ64936
ID ADQ64936 standard; cDNA; 3019 BP.
XX AC ADQ64936;
XX DT 07-OCT-2004 (first entry)
XX DE Novel human cDNA sequence #2097.
XX KW ss; gene; osteopathic; neuroprotective; nootropic; antiparkinsonian;
XX KW cytosstatic; gene therapy; diagnostic marker; morbid state; osteoporosis;
XX KW neurological disease; Alzheimer's disease; Parkinson's disease; dementia;
XX KW cancer.
XX OS Homo sapiens.
XX PN EP140981-A2.
XX PD 28-JUL-2004.
XX XX 21-JAN-2004; 2004EP-00001196.
XX XX 21-JAN-2003; 2003JP-00102206.
XX XX 03-MAY-2003; 2003JP-00131392.
XX XX (REAS-) RES ASSOC BIOTECHNOLOGY.
XX XX
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PI Isogai T, Sugiyama T, Otsuki T, Wakamatsu A, Sato H, Ishii S;
PI Yamamoto J, Isono Y, Nagai K, Irie R;
XX DR WPI: 2004-535376/52.
XX DR P-PSDB; ADQ67124.
XX PT Novel 2495 cDNA, useful for treating osteoporosis, neurological diseases,
XX PT Alzheimer's diseases, Parkinson's diseases, dementia and various cancers.
XX PS Claim 1; SEQ ID NO 2097; 2449pp; English.
XX CC The invention relates to 2495 novel polynucleotides (I) and their encoded
XX CC polypeptides, sequences hybridizing to these nucleotides, sequences
XX CC encoding partial polypeptides and sequences having 70% or 90% identity to
XX CC the nucleotide and protein sequences. The nucleotides and polypeptides
XX CC are useful as diagnostic markers or therapeutic target for the diseases
XX CC or morbid states. They are also useful for treating osteoporosis,
XX CC neurological diseases, Alzheimer's diseases, Parkinson's diseases,
XX CC dementia and various cancers. This sequence corresponds to a nucleotide
XX CC sequence of the invention.
XX SQ Sequence 3019 BP; 833 A; 670 C; 785 G; 731 T; 0 U; 0 Other;

Query Match 39.8%; Score 145.2; DB 12; Length 3019;
Best Local Similarity 71.8%; Pred. No. 4.7e-29;
Matches 221; Conservative 0; Mismatches 78; Indels 9; Gaps 2;

Qy 64 CTTTTCATTGTTTGTAGTCTTACTTATTTTATAGAGAAAGGCTTGTCTCGTCTATCTAGA 123
Db 1066 CTTGTTTTCTTTTCTTCTTTTCTTTTGGAGACAGGCTCTACTCTGTATCTCCAG 1125

Qy 124 TTGGAGTCGACGCGGTGTAATCATAGCTTACTGTAGTCTTTGAATTCCTGAGTTCAAGAGAT 183
Db 1126 CTGGAGTCGAGTGGCATGATCAGAGCTCACTGCAGCTTCTGCGGTTCAAGTGAT 1185

Qy 184 CTTTCTGCTCAGCTTCCAGGTCAGTGTAGACTGAGACTATATG-TGCTGTACCATGACACAGCTG 242
Db 1186 CTCCACCTCAGTCTCTGAGTAGCTGGGACTACAGGCATGTACCACCATGCTCAGCTA 1245

Qy 243 ATTTTAAATTTTTTTTGTAGAGATG-----AGTTGCCAGGCTGTCTTTGAATC 294
Db 1246 ATCTTAAATTTTTTTGTAGAGACAGGCTCTCACTTTGTGCCAGGCTGTCTTTGAATC 1305

Qy 295 CTGGCTGAGGTGATCTCTGCGTTGACCTCCCAAGTATCTTTAGACTACAGATGCACTC 354
Db 1306 CTGAGTTCAAGTGATTTCTCTGCTTGACCTCCCAAGTCTGGGATTACAGGTGTGAGC 1365

Qy 355 CACCACGC 362
Db 1366 CACCACAC 1373

RESULT 11
ABZ58995
ID ABZ58995 standard; DNA; 10735 BP.
XX AC ABZ58995;
XX DT 28-APR-2003 (first entry)
XX DE Human oncosuppressive gene (DRAGO) fragment.
XX KW Oncosuppressive; apoptotic; p53; p73; cytostatic; gene therapy; tumour;
XX KW DRAGO; human; gene; ds.
XX OS Homo sapiens.
XX PN WO2003006498-A2.
XX XX 23-JAN-2003.
XX XX 09-JUL-2002; 2002WO-EP007625.
XX XX
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PR 10-JUL-2001; 2001IT-MI001465.
XX
PA (NOVU-) NOVUSPHARMA SPA.
XX
XX Brogginini M, D'incalci M;
XX
XX WPI; 2003-221715/21.
XX
XX New oncosuppressive polypeptide, useful for preparing a medicament for
XX treating tumors.
XX
XX Claim 3; Page 34-37; 42pp; English.
XX
XX The invention relates to oncosuppressive polypeptides and encoding
XX polynucleotides. The oncosuppressive gene is involved in apoptotic
XX process and is regulated by p53 and p73. The oncosuppressive
XX polynucleotides are useful for preparing a medicament for treating
XX tumour. The present sequence represents a human oncosuppressive gene
XX (DRAGO) fragment, located upstream of the first exon
XX
XX Sequence 10735 BP; 2720 A; 2485 C; 2630 G; 2900 T; 0 U; 0 Other;
SQ
Query Match 39.7%; Score 145; DB 8; Length 10735;
Best Local Similarity 67.8%; Pred. No. 7.7e-29;
Matches 234; Conservative 0; Mismatches 105; Indels 6; Gaps 2;
QY 19 TTACTTAACCTCTGGGTTTAACCTTAACTTAACCCGTGTGTAATTTATTTTGTGATTTGTTTAA 78
Db ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 79 GTCTTACTTTATTTTGTAGAAAGGGTCTTGCTCCGTCATCTAGATTGGAGTCAGCGGT 138
Db ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 389 TTTTGTGGTTTTTTTGACACAGAGCTTTGCTGTGCACATATATATAATTTTAAATTTTAA 388
QY 139 GTAATCATAGCTTACTGTAGTCTTGAATTCCTCAGTTCAAGAGATCCTTCGCTCAGCT 198
Db ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 449 ACAATCACAGTTACTTATAGTCTGACCTCCGAGCTCAATTAATCTCCACCTCAGCC 508
QY 199 TCCAGGTAGCTGAGACTATATGTG-CTGTACTACATGCACAGCTGATTTTAAATTTT 257
Db ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 509 TCCTGAGTAGCTGGACCAACAGCGGTGTGCCACATCGCTGGCTAATTTTGTATTTT 568
QY 258 TTGTAGAG-----ATGAGGTGCCAGGCTGTGTTGAACTCCTGCGCTGAGTGATCCT 312
Db ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 569 GTAAAGAGGTCCTCACTCTGTGTCGCCAGGCTGGTCTCAAACTTCTGAGCTCAAGTGATCCT 628
QY 313 CTGCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACCTCCAC 357
Db ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 629 CTGCTTTGGCTTCCCAAGTGTGGGATTACAGGCGTGAGCCAC 673
RESULT 12
AAK88992/c
ID AAK88992 standard; DNA; 11216 BP.
XX
XX AAK88992;
AC
XX
XX 05-NOV-2001 (first entry)
DT
XX
XX Human digestive system antigen genomic sequence SEQ ID NO: 2568.
DE
XX
XX Human; digestive system antigen; gene therapy; cancer; appendicitis;
KW ulcerative colitis; infection; Hirschsprung's disease; chronic colitis;
KW digestive system disorder; Meckel's diverticulum; ds.
XX
XX Homo sapiens.
XX
XX WO20015314-A2.
XX
XX 02-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US001324.
XX
XX
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PR 07-JUL-2000; 2000US-0216647P.
PR 07-JUL-2000; 2000US-0216880P.
PR 11-JUL-2000; 2000US-0217487P.
PR 11-JUL-2000; 2000US-0217496P.
PR 14-JUL-2000; 2000US-0218290P.
PR 26-JUL-2000; 2000US-0220963P.
PR 26-JUL-2000; 2000US-0220964P.
PR 14-AUG-2000; 2000US-0224518P.
PR 14-AUG-2000; 2000US-0224519P.
PR 14-AUG-2000; 2000US-0225213P.
PR 14-AUG-2000; 2000US-0225214P.
PR 14-AUG-2000; 2000US-0225266P.
PR 14-AUG-2000; 2000US-0225267P.
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PR 14-AUG-2000; 2000US-0225447P.
PR 14-AUG-2000; 2000US-0225758P.
PR 14-AUG-2000; 2000US-0225759P.
PR 18-AUG-2000; 2000US-0226279P.
PR 22-AUG-2000; 2000US-0226681P.
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PR 01-SEP-2000; 2000US-0229287P.
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PR 06-SEP-2000; 2000US-0230437P.
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PR 08-SEP-2000; 2000US-0231242P.
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PR 08-SEP-2000; 2000US-0231413P.
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PR 08-SEP-2000; 2000US-0232080P.
PR 08-SEP-2000; 2000US-0232081P.
PR 12-SEP-2000; 2000US-0231968P.
PR 14-SEP-2000; 2000US-0232397P.
PR 14-SEP-2000; 2000US-0232398P.
PR 14-SEP-2000; 2000US-0232399P.
PR 14-SEP-2000; 2000US-0232400P.
PR 14-SEP-2000; 2000US-0232401P.
PR 14-SEP-2000; 2000US-0233063P.
PR 14-SEP-2000; 2000US-0233064P.
PR 14-SEP-2000; 2000US-0233065P.
PR 21-SEP-2000; 2000US-0234223P.
PR 21-SEP-2000; 2000US-0234274P.
PR 25-SEP-2000; 2000US-0234997P.
PR 25-SEP-2000; 2000US-0234998P.
PR 26-SEP-2000; 2000US-0235484P.
PR 27-SEP-2000; 2000US-0235834P.
PR 27-SEP-2000; 2000US-0235836P.
PR 29-SEP-2000; 2000US-0236327P.
PR 29-SEP-2000; 2000US-0236367P.
PR 29-SEP-2000; 2000US-0236368P.
PR 29-SEP-2000; 2000US-0236369P.
PR 29-SEP-2000; 2000US-0236370P.
PR 02-OCT-2000; 2000US-0236802P.
PR 02-OCT-2000; 2000US-0237037P.
PR 02-OCT-2000; 2000US-0237038P.
PR 02-OCT-2000; 2000US-0237039P.
PR 02-OCT-2000; 2000US-0237040P.
PR 13-OCT-2000; 2000US-0239935P.
PR 13-OCT-2000; 2000US-0239937P.
PR 20-OCT-2000; 2000US-0240960P.
PR 20-OCT-2000; 2000US-0241221P.
PR 20-OCT-2000; 2000US-0241785P.
PR 20-OCT-2000; 2000US-0241786P.
PR 20-OCT-2000; 2000US-0241787P.

PR 20-OCT-2000; 2000US-0241808P.
PR 20-OCT-2000; 2000US-0241809P.
PR 01-NOV-2000; 2000US-0244826P.
PR 01-NOV-2000; 2000US-0244617P.
PR 08-NOV-2000; 2000US-0246474P.
PR 08-NOV-2000; 2000US-0246475P.
PR 08-NOV-2000; 2000US-0246476P.
PR 08-NOV-2000; 2000US-0246477P.
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PR 08-NOV-2000; 2000US-0246523P.
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PR 08-NOV-2000; 2000US-0246532P.
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PR 08-NOV-2000; 2000US-0246611P.
PR 08-NOV-2000; 2000US-0246613P.
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PR 17-NOV-2000; 2000US-0249213P.
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PR 17-NOV-2000; 2000US-0249216P.
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PR 17-NOV-2000; 2000US-0249244P.
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PR 17-NOV-2000; 2000US-0249265P.
PR 17-NOV-2000; 2000US-0249297P.
PR 17-NOV-2000; 2000US-0249299P.
PR 17-NOV-2000; 2000US-0249300P.
PR 01-DEC-2000; 2000US-0250160P.
PR 01-DEC-2000; 2000US-0250391P.
PR 05-DEC-2000; 2000US-0251030P.
PR 05-DEC-2000; 2000US-0251988P.
PR 05-DEC-2000; 2000US-0256719P.
PR 06-DEC-2000; 2000US-0251479P.
PR 08-DEC-2000; 2000US-0251856P.
PR 08-DEC-2000; 2000US-0251868P.
PR 08-DEC-2000; 2000US-0251869P.
PR 08-DEC-2000; 2000US-0251989P.
PR 08-DEC-2000; 2000US-0251990P.
PR 11-DEC-2000; 2000US-0254097P.
PR 05-JAN-2001; 2001US-0259678P.

(HUMA-) HUMAN GENOME SCI INC.

PI Rosen CA, Barash SC, Ruben SM;

XX WPI; 2001-465567/50.

XX Isolated polypeptide for treating, preventing and/ or prognosing
PT disorders related to the colon including colon cancers and also for
PT testing and detection e.g. diagnosis.

XX Disclosure; SEQ ID NO 490; 562pp; English.

XX The present invention relates to the isolation of novel human colon
CC associated polypeptides (AAU22468-AAU22701), and the cDNA and genomic
CC sequences encoding for them. The sequences of the invention are useful in
CC the diagnosis, treatment, prevention and/or prognosis of disorders of the
CC colon including colon cancer, congenital abnormalities (e.g. atresia and
CC stenosis), bacterial and viral infections, inflammatory bowel disease
CC (IBD), neoplastic cell disorders (e.g. polyps and adenomas, intestinal
CC inflammatory disorders, colitis, colonic inflammation, diarrhoea and

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CC dysentery, malabsorption syndromes (e.g. lactose intolerance), intestinal
CC obstruction and sigmoid diseases. The polynucleotide sequences of the
CC instruction can also be used in gene therapy. AAS39582-AAS40060 represent
CC DNA sequences encoding for the novel human colon associated polypeptides
CC of the invention. Note: The sequence data for this patent did not form
CC part of the printed specification, but was obtained in electronic format
CC directly from WIPO at ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 11216 BP; 3857 A; 1984 C; 1812 G; 3563 T; 0 U; 0 Other;

Query Match 39.38; Score 143.6; DB 5; Length 11216;
Best Local Similarity 71.66; Pred. No. 1.9e-28;
Matches 202; Conservative 0; Mismatches 79; Indels 1; Gaps 1;

Qy 82 TTACTTTTATTTTAGAGAAAGGGTCTTGCTCCGTCATCTAGATTGGAGTGCAGCGGTGA 141
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 142 ATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGAGATCCTTCTGCCTCAGCTTCC 201
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 8180 ATCATTCCTCACTGCAGCCTTGAACTTCTGGGCTCAAGCAATCCTCCTGCCTCAGATTCC 8121
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 202 CAGGTAGCTGAGACTATATGTC-TGCTACCATGCACAGCTGATTTTAAATTTTTTGG 260
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 8120 CAGGTAGCTGGGACTATAGGCACATGCCACCATCCCGAGCTAATTTTGTAGAGACAG 8061
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 261 TAGAGATGGAGTTGCCAGGCTGTCTTTGAACCTCTGGCTGAGGTGATCCTCTCGGTT 320
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 8060 GTCTCACTGTGTGCCAGGCTGTCTCGAACTTCTGGGCTCAAGCTGTCTCTCGCTT 8001
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 321 GACCTCCCAAGTATCTTTAGACTACAGATGCACTCCACACGC 362
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 8000 AACCTCCAAATACTGGGATTACAGACATGAGCCACACAC 7959
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

RESULT 14
ADB32553/c
ID ADB32553 standard; DNA; 11216 BP.
XX
AC ADB32553;
XX
DT 04-DEC-2003 (first entry)
XX
DE Human novel colon related polypeptide DNA SEQ ID NO 490.
XX
KW gene therapy; cancer; liver disorder; hepatitis; neural disorder;
KW Alzheimer's disease; human; colon; ds.
XX
OS Homo sapiens.
XX
PN US2003050231-A1.
XX
PD 13-MAR-2003.
XX
XX 17-JAN-2001; 2001US-00764872.
XX
XX 31-JAN-2000; 2000US-0179065P.
XX 04-FEB-2000; 2000US-0180628P.
XX 24-FEB-2000; 2000US-0184664P.
XX 02-MAR-2000; 2000US-0186350P.
XX 16-MAR-2000; 2000US-0189874P.
XX 17-MAR-2000; 2000US-0190076P.
XX 18-APR-2000; 2000US-0198123P.
XX 19-MAY-2000; 2000US-0205515P.
XX 07-JUN-2000; 2000US-0209467P.
XX 28-JUN-2000; 2000US-0214886P.
XX 30-JUN-2000; 2000US-0215135P.
XX 07-JUL-2000; 2000US-0216647P.
XX 07-JUL-2000; 2000US-0216880P.
XX 11-JUL-2000; 2000US-0217487P.
XX 11-JUL-2000; 2000US-0217496P.
XX 14-JUL-2000; 2000US-0218290P.
XX 26-JUL-2000; 2000US-0220963P.
PR 26-JUL-2000; 2000US-0220964P.
PR 14-AUG-2000; 2000US-0224518P.
PR 14-AUG-2000; 2000US-0224519P.
PR 14-AUG-2000; 2000US-0225213P.
PR 14-AUG-2000; 2000US-0225214P.
PR 14-AUG-2000; 2000US-0225266P.
PR 14-AUG-2000; 2000US-0225267P.
PR 14-AUG-2000; 2000US-0225268P.
PR 14-AUG-2000; 2000US-0225370P.
PR 14-AUG-2000; 2000US-0225447P.
PR 14-AUG-2000; 2000US-0225757P.
PR 14-AUG-2000; 2000US-0225758P.
PR 14-AUG-2000; 2000US-0225759P.
PR 18-AUG-2000; 2000US-0226279P.
PR 22-AUG-2000; 2000US-0226681P.
PR 22-AUG-2000; 2000US-0226686P.
PR 22-AUG-2000; 2000US-0227182P.
PR 23-AUG-2000; 2000US-0227009P.
PR 30-AUG-2000; 2000US-0228924P.
PR 01-SEP-2000; 2000US-0229287P.
PR 01-SEP-2000; 2000US-0229343P.
PR 01-SEP-2000; 2000US-0229344P.
PR 01-SEP-2000; 2000US-0229345P.
PR 05-SEP-2000; 2000US-0229509P.
PR 05-SEP-2000; 2000US-0229513P.
PR 06-SEP-2000; 2000US-0230437P.
PR 06-SEP-2000; 2000US-0230438P.
PR 08-SEP-2000; 2000US-0231242P.
PR 08-SEP-2000; 2000US-0231243P.
PR 08-SEP-2000; 2000US-0231244P.
PR 08-SEP-2000; 2000US-0231413P.
PR 08-SEP-2000; 2000US-0231414P.
PR 08-SEP-2000; 2000US-0232080P.
PR 08-SEP-2000; 2000US-0232081P.
PR 12-SEP-2000; 2000US-0231968P.
PR 14-SEP-2000; 2000US-0232397P.
PR 14-SEP-2000; 2000US-0232398P.
PR 14-SEP-2000; 2000US-0232399P.
PR 14-SEP-2000; 2000US-0232400P.
PR 14-SEP-2000; 2000US-0232401P.
PR 14-SEP-2000; 2000US-0233063P.
PR 14-SEP-2000; 2000US-0233064P.
PR 21-SEP-2000; 2000US-0234223P.
PR 21-SEP-2000; 2000US-0234274P.
PR 25-SEP-2000; 2000US-0234997P.
PR 25-SEP-2000; 2000US-0234998P.
PR 26-SEP-2000; 2000US-0235483P.
PR 27-SEP-2000; 2000US-0235834P.
PR 27-SEP-2000; 2000US-0235836P.
PR 29-SEP-2000; 2000US-0236327P.
PR 29-SEP-2000; 2000US-0236367P.
PR 29-SEP-2000; 2000US-0236368P.
PR 29-SEP-2000; 2000US-0236369P.
PR 29-SEP-2000; 2000US-0236370P.
PR 02-OCT-2000; 2000US-0237037P.
PR 02-OCT-2000; 2000US-0237038P.
PR 02-OCT-2000; 2000US-0237039P.
PR 13-OCT-2000; 2000US-0239935P.
PR 13-OCT-2000; 2000US-0239937P.
PR 20-OCT-2000; 2000US-0240960P.
PR 20-OCT-2000; 2000US-0241221P.
PR 20-OCT-2000; 2000US-0241785P.
PR 20-OCT-2000; 2000US-0241786P.
PR 20-OCT-2000; 2000US-0241787P.
PR 20-OCT-2000; 2000US-0241808P.
PR 20-OCT-2000; 2000US-0241809P.
PR 20-OCT-2000; 2000US-0241826P.
PR 01-NOV-2000; 2000US-0244617P.
PR 08-NOV-2000; 2000US-0246474P.
PR 08-NOV-2000; 2000US-0246475P.
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OM nucleic - nucleic search, using sw model

Run on: August 30, 2005, 03:39:08 ; Search time 136 Seconds
(without alignments)
4391.478 Million cell updates/sec

Title: US-08-731-499-9_COPY_10001_10365

Perfect score: 365

Sequence: 1 TTTGTGGTCTCCAGGCTT.....GATGCACTCCACGCGTTG 365

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents NA.*

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- 2: /cgn2_6/ptodata/1/ina/5B_COMB.seq.*
- 3: /cgn2_6/ptodata/1/ina/6A_COMB.seq.*
- 4: /cgn2_6/ptodata/1/ina/6B_COMB.seq.*
- 5: /cgn2_6/ptodata/1/ina/PCTUS_COMB.seq.*
- 6: /cgn2_6/ptodata/1/ina/backfilesi.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	365	100.0	10365	4	US-08-892-695-9
2	365	100.0	20022	4	US-09-949-016-12604
3	365	100.0	20023	4	US-09-949-016-16004
4	149.2	40.9	38538	4	US-09-949-016-13150
5	146.8	40.2	197131	4	US-09-949-016-12675
6	146.8	40.2	197132	4	US-09-949-016-17170
c 7	145.2	39.8	22010	4	US-09-949-016-15960
8	144.8	39.7	601	4	US-09-949-016-51283
9	144.8	39.7	73295	4	US-09-949-016-15151
10	144.8	39.7	121427	4	US-09-949-016-11950
c 11	144.8	39.7	121433	4	US-09-949-016-13230
c 12	144.8	39.6	25736	4	US-09-949-016-15090
c 13	144.4	39.6	25755	4	US-09-949-016-12351
c 14	144.4	39.6	25755	4	US-09-949-016-12351
c 15	144.2	39.5	601	4	US-09-949-016-121870
c 16	143	39.2	33392	4	US-09-949-016-15172
17	142.4	39.0	24665	4	US-09-949-016-17134
18	142.2	39.0	36759	4	US-09-949-016-12216
19	142.2	39.0	36760	4	US-09-949-016-14021
c 20	142	38.9	601	4	US-09-949-016-157469
c 21	142	38.9	601	4	US-09-949-016-157576
c 22	142	38.9	601	4	US-09-949-016-17521
c 23	141	38.6	87734	4	US-09-949-016-15341
c 24	141	38.6	235452	4	US-09-949-016-13675
c 25	140.8	38.6	601	4	US-09-949-016-141029
c 26	140.8	38.6	601	4	US-09-949-016-141030
c 27	140.8	38.6	601	4	US-09-949-016-157470

ALIGNMENTS

RESULT 1

US-08-892-695-9

; Sequence 9, Application US/08892695A

; Patent No. 6808878

; GENERAL INFORMATION:

; APPLICANT: Gray, Joe W

; APPLICANT: Collins, Collin

; APPLICANT: Hwang, Soo In

; APPLICANT: Godfrey, Tony

; APPLICANT: Kowel, David

; APPLICANT: Rommens, Johanna

; TITLE OF INVENTION: GENES FROM THE 20Q13 AMPLICON AND THEIR USES

; FILE REFERENCE: 2500.124US3

; CURRENT APPLICATION NUMBER: US/08/892,695A

; CURRENT FILING DATE: 1997-07-15

; EARLIER APPLICATION NUMBER: 08/785,532

; EARLIER FILING DATE: 1997-01-17

; EARLIER APPLICATION NUMBER: 08/731,499

; EARLIER FILING DATE: 1996-10-16

; EARLIER APPLICATION NUMBER: 08/680,395

; EARLIER FILING DATE: 1996-07-15

; NUMBER OF SEQ ID NOS: 59

; SOFTWARE: PatentIn Ver. 2.0

; SEQ ID NO 9

; LENGTH: 10365

; TYPE: DNA

; ORGANISM: Artificial Sequence

; FEATURE:

; OTHER INFORMATION: Description of Artificial Sequence: Genomic

; OTHER INFORMATION: Sequence encoding ZABC1

; Patent No. 6808878

US-08-892-695-9

Query Match 100.0%; Score 365; DB 4; Length 10365;

Best Local Similarity 100.0%; Pred. No. 1.1e-100;

Matches 365; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TTTGTGGTCTCCAGGCTTAACTCTGTGGGTTAACTCTTAACCCCTGTGATTTT 60

Db 10001 TTTGTGGTCTCCAGGCTTAACTCTGTGGGTTAACTCTTAACCCCTGTGATTTT 10060

Qy 61 ATTCTTTGATTTTGTAGTCTTACTTTATTTTAGAAGAGGCTCTTGTCCGTCATCT 120

Db 10061 ATTCTTTGATTTTGTAGTCTTACTTTATTTTAGAAGAGGCTCTTGTCCGTCATCT 10120

Qy 121 AGATTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGA 180

Db 10121 AGATTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGA 10180

Qy 181 GATCCTTCTGCTCAGCTTCCAGGTAGCTGAGACTATATGTGCTGCTACCATCAGC 240

Sequence 157577, A

Sequence 16137, A

Sequence 16138, A

Sequence 121481, A

Sequence 157471, A

Sequence 157578, A

Sequence 15131, A

Sequence 14606, A

Sequence 89655, A

Sequence 89666, A

Sequence 89677, A

Sequence 89688, A

Sequence 15086, A

Sequence 17390, A

Sequence 17391, A

Sequence 6, Appli

Sequence 1, Appli

Sequence 89656, A

Db 10181 GATCCTTCTGCCCTCAGCTCCAGGTAGCTGAGACTATATGTGCTGTACCATGCACAGC 10240
Qy 241 TGATTTTAAATTTTTTTGTAGAGATGGAGTTGCCAGGCTGGTCTTTGAACCTCTGGCC 300
Db 10241 TGATTTTAAATTTTTTTGTAGAGATGGAGTTGCCAGGCTGGTCTTTGAACCTCTGGCC 10300
Qy 301 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACCTCCACCAC 360
Db 10301 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACCTCCACCAC 10360
Qy 361 GCTTG 365
Db 10361 GCTTG 10365

RESULT 2

US-09-949-016-12604
; Sequence 12604, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12604
; LENGTH: 20022
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-12604

Query Match 100.0%; Score 365; DB 4; Length 20022;
Best Local Similarity 100.0%; Pred. No. 1.5e-100;
Matches 365; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 TTTGTGCTCTCCAAGGCTTACTTAACCTCTGTGGGTTTAACTCTTAACCCCTGTGATTTT 60
Db 9531 TTTGTGCTCTCCAAGGCTTACTTAACCTCTGTGGGTTTAACTCTTAACCCCTGTGATTTT 9590
Qy 61 ATCTTTTGAATTTGTTAGTCTTACTTTATTTTAGAAGAGGCTTGTCTCCGTCACTCT 120
Db 9591 ATCTTTTGAATTTGTTAGTCTTACTTTATTTTAGAAGAGGCTTGTCTCCGTCACTCT 9650
Qy 121 AGATTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTTCCCTGAGTTCAAGA 180
Db 9651 AGATTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTTCCCTGAGTTCAAGA 9710
Qy 181 GATCCTTCTGCCCTCAGCTTCCCAGGTAGCTGAGACTATATGTGCTGTACCATGCACAGC 240
Db 9711 GATCCTTCTGCCCTCAGCTTCCCAGGTAGCTGAGACTATATGTGCTGTACCATGCACAGC 9770
Qy 241 TGATTTTAAATTTTTTTGTAGAGATGGAGTTGCCAGGCTGGTCTTGAACCTCTGGCC 300
Db 9771 TGATTTTAAATTTTTTTGTAGAGATGGAGTTGCCAGGCTGGTCTTGAACCTCTGGCC 9830
Qy 301 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACCTCCACCAC 360
Db 9831 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACCTCCACCAC 9890
Qy 361 GCTTG 365
Db 9891 GCTTG 9895

RESULT 3

US-09-949-016-16004
; Sequence 16004, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16004
; LENGTH: 20023
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-16004

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Best Local Similarity 100.0%; Pred. No. 1.5e-100;
Matches 365; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 TTTGTGCTCTCCAAGGCTTACTTAACCTCTGTGGGTTTAACTCTTAACCCCTGTGATTTT 60
Db 9531 TTTGTGCTCTCCAAGGCTTACTTAACCTCTGTGGGTTTAACTCTTAACCCCTGTGATTTT 9590
Qy 61 ATCTTTTGAATTTGTTAGTCTTACTTTATTTTAGAAGAGGCTTGTCTCCGTCACTCT 120
Db 9591 ATCTTTTGAATTTGTTAGTCTTACTTTATTTTAGAAGAGGCTTGTCTCCGTCACTCT 9650
Qy 121 AGATTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTTCCCTGAGTTCAAGA 180
Db 9651 AGATTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTTCCCTGAGTTCAAGA 9710
Qy 181 GATCCTTCTGCCCTCAGCTTCCCAGGTAGCTGAGACTATATGTGCTGTACCATGCACAGC 240
Db 9711 GATCCTTCTGCCCTCAGCTTCCCAGGTAGCTGAGACTATATGTGCTGTACCATGCACAGC 9770
Qy 241 TGATTTTAAATTTTTTTGTAGAGATGGAGTTGCCAGGCTGGTCTTGAACCTCTGGCC 300
Db 9771 TGATTTTAAATTTTTTTGTAGAGATGGAGTTGCCAGGCTGGTCTTGAACCTCTGGCC 9830
Qy 301 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACCTCCACCAC 360
Db 9831 TGAGGTGATCCTCTCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACCTCCACCAC 9890
Qy 361 GCTTG 365
Db 9891 GCTTG 9895

RESULT 4

US-09-949-016-13150
; Sequence 13150, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768

Qy	40	ACTCTTAACCCCTGTGTAATTTTATCTCTTTGATTTGTTTAGTCTTACTTTATTTTGTAGAGA	99
Db	134568	ACTCATTTGTAATCTGTGTGTGTGTTTTTTTTTTGTTTTTGTGTTTTTTTTTTTGTAGAGA	134627
Qy	100	AAGGGTCTTGCTCCGTCATCTAGATTTGGAGTGGAGCGGCTGTAATCATAGCTTACTGTAGT	159
Db	134628	CAGAGTCTCACTCTGTCAACCAGGCTGGAGTGGAGTGGTGATCTCAGCTCACTGCAAC	134687
Qy	160	CTTGAATTCCTGAGTTCAAGAGATCCCTTTCGCTCAGCTTCCCAAGGTAGCTGAGACTATA	219
Db	134688	CTCCACCTCTCGGGTTCAAGCAATTCCTCTGCCTTAGCTCCCAAGTAGCTGGGACTATA	134747
Qy	220	TGTGTC-TGCTACCATGCACAGCTGATTTTTTAAATTTTTTTTGTAGATGG-----A	270
Db	134748	GGCGCAGCCACCAACGCCAGCTAAATTTTTTGTATTTTTTAGTAGACAGGGTTTTTACCAC	134807
Qy	271	GTTGCCCGAGCTGTGCTTTGAACCTCTGGCCTGAGGTGATCTCTCGCTTGACCTCCCAA	330
Db	134808	GTTGGCCAGACCGGTCTTTGAACCTCTGACCTCAGGTGATCTCCCACTCGGCTCCCAA	134867
Qy	331	GTATCTTTAGACTACAGATGCACCTCCACCAGC	362
Db	134868	GGTGCTGGATTACAGGTGTGAGCCACCGTGC	134899
RESULT 6			
US-09-949-016-17170			
; Sequence 17170, Application US/09949016			
; Patent No. 6812339			
; GENERAL INFORMATION:			
; APPLICANT: VENTER, J. Craig et al.			
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED			
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF			
; FILE REFERENCE: CL001307			
; CURRENT APPLICATION NUMBER: US/09/949,016			
; CURRENT FILING DATE: 2000-04-14			
; PRIOR APPLICATION NUMBER: 60/241,755			
; PRIOR FILING DATE: 2000-10-20			
; PRIOR APPLICATION NUMBER: 60/237,768			
; PRIOR FILING DATE: 2000-10-03			
; PRIOR APPLICATION NUMBER: 60/231,498			
; PRIOR FILING DATE: 2000-09-08			
; NUMBER OF SEQ ID NOS: 207012			
; SOFTWARE: FastSeq for Windows Version 4.0			
; SEQ ID NO 17170			
; LENGTH: 197132			
; TYPE: DNA			
; ORGANISM: Human			
; FEATURE:			
; NAME/KEY: misc_feature			
; LOCATION: (1)..(197132)			
; OTHER INFORMATION: n = A, T, C or G			
US-09-949-016-17170			
Query Match 40.2%; Score 146.8; DB 4; Length 197132;			
Best Local Similarity 69.6%; Pred. No. 9.3e-34;			
Matches 231; Conservative 0; Mismatches 92; Indels 9; Gaps 2;			
Qy	40	ACTCTTAACCCCTGTGTAATTTTATCTCTTTGATTTGTTTAGTCTTACTTTATTTTGTAGAGA	99
Db	134568	ACTCATTTGTAATCTGTGTGTGTGTTTTTTTTTTGTTTTTGTGTTTTTTTTTTTGTAGAGA	134627
Qy	100	AAGGGTCTTGCTCCGTCATCTAGATTTGGAGTGGAGCGGCTGTAATCATAGCTTACTGTAGT	159
Db	134628	CAGAGTCTCACTCTGTCAACCAGGCTGGAGTGGAGTGGTGATCTCAGCTCACTGCAAC	134687
Qy	160	CTTGAATTCCTGAGTTCAAGAGATCCCTTTCGCTCAGCTTCCCAAGGTAGCTGAGACTATA	219
Db	134688	CTCCACCTCTCGGGTTCAAGCAATTCCTCTGCCTTAGCTCCCAAGTAGCTGGGACTATA	134747
Qy	220	TGTGTC-TGCTACCATGCACAGCTGATTTTTTAAATTTTTTTTGTAGATGG-----A	270
Db	134748	GGCGCAGCCACCAACGCCAGCTAAATTTTTTGTATTTTTTAGTAGACAGGGTTTTTACCAC	134807
Qy	271	GTTGCCCGAGCTGTGCTTTGAACCTCTGGCCTGAGGTGATCTCTCGCTTGACCTCCCAA	330
Db	134808	GTTGGCCAGACCGGTCTTTGAACCTCTGACCTCAGGTGATCTCCCACTCGGCTCCCAA	134867
Qy	331	GTATCTTTAGACTACAGATGCACCTCCACCAGC	362
Db	134868	GGTGCTGGATTACAGGTGTGAGCCACCGTGC	134899

; PRIOR FILING DATE: 2000-10-03
 ; PRIOR APPLICATION NUMBER: 60/231,498
 ; PRIOR FILING DATE: 2000-09-08
 ; NUMBER OF SEQ ID NOS: 207012
 ; SOFTWARE: FastSeq for Windows Version 4.0
 ; SEQ ID NO 13150
 ; LENGTH: 38538
 ; TYPE: DNA
 ; ORGANISM: Human
 ; FEATURE:
 ; NAME/KEY: misc feature
 ; LOCATION: (1)...(38538)
 ; OTHER INFORMATION: n = A,T,C or G
 US-09-949-016-13150

Query Match 40.9%; Score 149.2; DB 4; Length 38538;
 Best Local Similarity 75.4%; Pred. No. 8.1e-35;
 Matches 214; Conservative 0; Mismatches 63; Indels 7; Gaps 2;

Qy	83	TACTTTATTTT	TAGGAAGGGTCTTGCTCCGTCACTAGATGCAAGTGCAGCGGTAA	142
Db	5869	TAATTTGTTTT	TAAGAGACAGGGTCTCACTCTGTCAATCAGATGAAGTGCAGTGAGTGA	5928
Qy	143	TCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGAGATCCTTCTGCCTCAGCTTCCC	202	
Db	5929	TCATAGCTCACTGCAGCTAGACTCTCTGGGCTCAAGTGATCCTCTGCCTCAGCTCTC	5988	
Qy	203	AGGTAGCTGAGACTATATGTGC-TGCTACCATGCACAGCTGAATTTTAAATTTTTTGT	261	
Db	5989	CAGTAGCTGAGACTATAGGCACATGCCACCACATGCCAGCTAATTTTAAACTTTTTTGT	6048	
Qy	262	AGAGATGGAGTTGCCAGGCTGGCTTGAACCTCTGGCTGAGGTGATCCTCTGGGTG	321	
Db	6049	AAGGA-----TGCTGAGGCTGGTCTTGAACCTCTGGCATCAAGCAATCCTCCCGCTTA	6102	
Qy	322	ACCTCCCAAGTATCTTAGACTACAGATGCATCTCCACACAGCTTG	365	
Db	6103	GCCTCCCAATCTTGGGATTAGAGCGTGAGCCACCATGCTTG	6146	

RESULT 5
 US-09-949-016-12675
 ; Sequence 12675, Application US/09949016
 ; Patent No. 6812339
 ; GENERAL INFORMATION:
 ; APPLICANT: VENTER, J. Craig et al.
 ; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
 ; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
 ; FILE REFERENCE: CL001307
 ; CURRENT APPLICATION NUMBER: US/09/949,016
 ; CURRENT FILING DATE: 2000-04-14
 ; PRIOR APPLICATION NUMBER: 60/241,755
 ; PRIOR FILING DATE: 2000-10-20
 ; PRIOR APPLICATION NUMBER: 60/237,768
 ; PRIOR FILING DATE: 2000-10-03
 ; PRIOR APPLICATION NUMBER: 60/231,498
 ; PRIOR FILING DATE: 2000-09-08
 ; NUMBER OF SEQ ID NOS: 207012
 ; SOFTWARE: FastSeq for Windows Version 4.0
 ; SEQ ID NO 12675
 ; LENGTH: 197131
 ; TYPE: DNA
 ; ORGANISM: Human
 ; FEATURE:
 ; NAME/KEY: misc feature
 ; LOCATION: (1)...(197131)
 ; OTHER INFORMATION: n = A,T,C or G
 US-09-949-016-12675

Query Match 40.2%; Score 146.8; DB 4; Length 197131;
 Best Local Similarity 69.6%; Pred. No. 9.3e-34;
 Matches 231; Conservative 0; Mismatches 92; Indels 9; Gaps 2;


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Query Match          39.7%; Score 144.8; DB 4; Length 52971;
Best Local Similarity 70.2%; Pred. No. 2e-33;
Matches 212; Conservative 0; Mismatches 82; Indels 8; Gaps 1;

Qy 72 TTTTGTAGCTTACTTATTTTATAGAGAAAGGCTTGTCTCCGTCATCTAGATTGGAGTG 131
Db 21346 TTTTGTAGCTTACTTATTTTATAGAGAAAGGCTTGTCTCCGTCATCTAGATTGGAGTG 131
Qy 132 CAGCGGTGTAATCATAGCTTACTAGTCTTGAATTCCTGAGTTCAAGAGATCCTCTGC 191
Db 21406 CAGCAGGCAATCTTAGTCTACTGAGCTCCACCTCTCTGGGTTCAACAAATTCCTCTGC 21465
Qy 192 CTCAGCTTCCAGGAGTAGCTAGACTATATGTGCTGTACCATGCACAGCTGATTTTAAA 251
Db 21466 CTCAGCTTCCAGGAGTAGCTAGACTATATGTGCTGTACCATGCACAGCTGATTTTAAA 251
Qy 252 TTTTGTAGAGATGG-----AGTGTCCAGGCTGTGCTTGAACCTCTGGCTGA 303
Db 21526 TTTTGTAGAGACAGAGGTTTGTGCATGTTTGTCCAGGCTGTGCTTGAACCTCTGACCTCA 21585
Qy 304 GGTGATCCTCTGCTGTTGACCTCCCAAGTATCTTTAGACTACAGATGCACCTCCACGCT 363
Db 21586 AGTATCCACCTGTCTCCACCTCCCAAGTATCTTTAGACTACAGATGCACCTCCACGCT 363
Qy 364 TG 365
Db 21646 TG 21647

RESULT 10
US-09-949-016-15151
; Sequence 15151, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15151
; LENGTH: 73295
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)...(73295)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-15151

Query Match          39.7%; Score 144.8; DB 4; Length 73295;
Best Local Similarity 72.8%; Pred. No. 2.4e-33;
Matches 217; Conservative 0; Mismatches 72; Indels 9; Gaps 2;

Qy 75 TTTAGTCTTACTTATTTTATAGAGAAAGGCTTGTCTCCGTCATCTAGATTGGAGTGAC 134
Db 49031 TTGTTTTTTTTTTTTTTTTTGTAGACAGGGTCTTACTCTGTCAACCCAGCTGGAGTGAC 49090
Qy 135 CGGTGTAATCATAGCTTACTAGTCTTGAATTCCTGAGTTCAAGAGATCCTCTGCTC 194
Db 49091 TGGCACAATCATAGCTTACTAGCTTACCGCCAGGCTCAAGACATCTCCACCCC 49150
Qy 195 AGCTTCCAGGAGTAGCTAGACTATATGTG-CTGCTACCATGCACAGCTGATTTTAAATT 253
Db 195 AGCTTCCAGGAGTAGCTAGACTATATGTG-CTGCTACCATGCACAGCTGATTTTAAATT 253
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Db 49151 AGCTTCCAAAGTAGCTGGGACTACAGGTGTGTGCCACCATGCCAGCTAATTTTGTATT 49210
Qy 254 TTTTGTAGAGATGG-----AGTGTCCAGGCTGGTCTTGAACCTCTGGCTGAGG 305
Db 49211 TTTTGTAGAAACAAGGTTTCCCATGTTGCCAGGCTGGTCTTGAACCTCTGGGACAGG 49270
Qy 306 TGATCTCTCTGCGTTGACCTCCCAAGPATCTTATAGACTACAGATGCACCTCCACCGCT 363
Db 49271 GGATCTGCTGCTGGGCTCCCAAGAGTGTGGGATTACAGATGGGAGCCACCACT 49328

RESULT 11
US-09-949-016-11950/c
; Sequence 11950, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 11950
; LENGTH: 121427
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-11950

Query Match          39.7%; Score 144.8; DB 4; Length 121427;
Best Local Similarity 67.2%; Pred. No. 3e-33;
Matches 238; Conservative 0; Mismatches 107; Indels 9; Gaps 2;

Qy 20 TACTTAACCTCTGCGGTTTAACCTCTTAACCTCTGATATTTTATTTCTTTGATTTGTTAG 79
Db 33363 TACTAAATACTACTCTTTCTTATGCAAAACTAGTATTTCAAAATATACAGATTTCTTTTC 33304
Qy 80 TCTTACTTTTATTTTATAGAGAAAGGCTCTGCTCGTCATCTAGATTGCGAGTGCAGCGTG 139
Db 33303 TTTTGTAGACAGAGTCTCACTGTGTCATCGAGCTGAGTGCAGTGGCG 33244
Qy 140 TAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAAGAGATCCTTCTGCTCAGCTT 199
Db 33243 TGATCTTGGCTCACTGCAACCTTTGCTCTCTGGGTTCAAGTGATTTCTCTGCTCAGCTT 33184
Qy 200 CCAGGTAGCTGAGACTATATGTG-CTGCTACCATGCACAGCTGATTTTAAATTTTTT 258
Db 33183 CCCAGTAGCTGAGATTACAGGTGCCACCATGCCCGCTAAATTTTGTATTTTATG 33124
Qy 259 TGTAGAGATGG-----AGTTGCCAGGCTGTGCTTGAATCTCTGCTGAGTGCATC 310
Db 33123 TAAAGACGGGTTTTCACCATGTTGGCCAGGCTGGTCTTGAACCTCTCACTCAATGATC 33064
Qy 311 CTCTGCTGTTGACCTCCCAAGTATCTTATAGACTACAGATGCACCTCCACGCTT 364
Db 33063 CGCCCGCTCGGCTCCCAAGTGTGGGATTACAGGATTAACAGCATGMAACCAACGCTT 33010

RESULT 12
US-09-949-016-13230/c
; Sequence 13230, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
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; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13230
; LENGTH: 121433
; TYPE: DNA
; ORGANISM: Human
; US-09-949-016-13230

Query Match      39.7%; Score 144.8; DB 4; Length 121433;
Best Local Similarity 67.2%; Pred. No. 3e-33;
Matches 238; Conservative 0; Mismatches 107; Indels 9; Gaps 2;

Qy 20 TACTTAACCTCTGCGGTTTAACTCTTAACCCCTGCTATTTTATTTCTTTTGGATTGTTTAG 79
Db 33362 TACTAAATACCTCTTTCTTATGCAAAACTAGTATTTTCAAAAATATCAGATTTCTTTTC 33303

Qy 80 TCCTACTTTATTTTAGAAGAGGCTCTGCTCGTCACTAGATTTGGAGTGCAGCGGTG 139
Db 33302 TTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTT 33243

Qy 140 TAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAGAGATCCTTCTGCTCAGCTT 199
Db 33242 TGAATCTTGGCTCACTGCACTTTGCTCTCTGCTTCAAGTATTTCTTCTGCTCAGCTT 33183

Qy 200 CCCAGGTAGCTGACATATATGTG-CTGCTACCATGCACAGCTGATTTTAAATTTTTTT 258
Db 33182 CCCAGGTAGCTGAGATTACAGGTGCCCCACCATGCCGGCTAAATTTTGTATTTTAG 33123

Qy 259 TGTAGAGATGG-----AGTTGCCAGGCTGGTCTTGAACCTCGCTGAGGTGATC 310
Db 33122 TAAAGACGGGGTTTTCACCATGTTGGCCAGGCTGGTCTTGAACCTCCTGACCTCAAGTGATC 33063

Qy 311 CTCTCGGTTGACCTCCCAAGTATCTTTAGACTACAGATGCACCTCCACACGCTT 364
Db 33062 CGCCCGCTCGGCTCCCAAGTCTGGGATTACAGCATGACCAACACGCTT 33009

RESULT 13
US-09-949-016-15090/c
; Sequence 15090 Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15090
; LENGTH: 25736
; TYPE: DNA
; ORGANISM: Human
; US-09-949-016-15090

Query Match      39.6%; Score 144.4; DB 4; Length 25736;
Best Local Similarity 70.4%; Pred. No. 1.9e-33;
Matches 207; Conservative 0; Mismatches 86; Indels 1; Gaps 1;

Qy 54 GTATTTTATTTCTTTGATTGTTTGTAGTCTTACTTTATTTTATTTAGAGAAAGGCTTTGCTCC 113
Db 20931 GTCTACTCTACATATGATATTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTT 20872

Qy 114 GTCATCTAGATTGGAGTGCAGCGGTGTAATCATAGTCTTACTGTAGTCTTGAATTCCTGAG 173
Db 20871 GTCGCCAGGCTGAAGTGCAGTGGTGCGATCTTGGCTCACTGCAACCTTTGCTTCCAGG 20812

Qy 174 TTCAAGAGATCCTTCTGCTCAGCTTCCAGAGTGCAGACTATATGTG-CTGCTACCA 232
Db 20811 TTCAAGGATTTCTCTCTGCTCAGCTTCCAGTGCAGGCTTACAGGCGTGTGCCACCA 20752

Qy 233 TGCACAGCTGATTTTAAATTTTGTAGAGATGGAGTTGCCAGGCTGGTCTTGAAC 292
Db 20751 TGCCAGCTAAATTTTCTGTATTTTGTAGAGATGAGGTGGCCAGGCTGGTCTTGAAC 20692

Qy 293 TCCTGCGCTGAGGTGATCTCTGCTGCTGACCTCCCAAGTATCTTAGACTACAG 346
Db 20691 TCCTGACCTCAAGTATCCACCTCGGCTCGGCTCCCAAGCGCTAGGATTACAG 20638

RESULT 14
US-09-949-016-12351/c
; Sequence 12351 Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12351
; LENGTH: 25755
; TYPE: DNA
; ORGANISM: Human
; US-09-949-016-12351

Query Match      39.6%; Score 144.4; DB 4; Length 25755;
Best Local Similarity 70.4%; Pred. No. 1.9e-33;
Matches 207; Conservative 0; Mismatches 86; Indels 1; Gaps 1;

Qy 54 GTATTTTATTTCTTTGATTGTTTGTAGTCTTACTTTATTTTATTTAGAGAAAGGCTTTGCTCC 113
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Db 20890 GTCGCCAGGCTGAAGTGCAGTGGTGCGATCTTGGCTCACTGCAACCTTTGCTTCCAGG 20831

Qy 174 TTCAAGAGATCCTTCTGCTCAGCTTCCAGAGTGCAGACTATATGTG-CTGCTACCA 232
Db 20830 TTCAAGGATTTCTCTCTGCTCAGCTTCCAGTGCAGGCTTACAGGCGTGTGCCACCA 20771

Qy 233 TGCACAGCTGATTTTAAATTTTGTAGAGATGGAGTTGCCAGGCTGGTCTTGAAC 292
Db 20770 TGCCAGCTAAATTTTCTGTATTTTGTAGAGATGAGGTGGCCAGGCTGGTCTTGAAC 20711

Qy 293 TCCTGCGCTGAGGTGATCTCTGCTGCTGACCTCCCAAGTATCTTAGACTACAG 346
Db 20710 TCCTGACCTCAAGTATCCACCTCGGCTCGGCTCCCAAGCGCTAGGATTACAG 20657
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RESULT 15

US-09-949-016-121870/c
; Sequence 121870, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 121870
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-121870

Query Match 39.5%; Score 144.2; DB 4; Length 601;
Best Local Similarity 73.5%; Pred. No. 3.9e-34;
Matches 214; Conservative 0; Mismatches 58; Indels 9; Gaps 2;

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Qy	142	ATCATAGCTTACT	TAGTCTTGATTCCTGAGTTCAGAGATCCTTCTGCCTCAGCTTC	201
Db	540	ATCATAGCTCACT	GTACGCTTGACCCGCGCTCAAGACATCTCCACCCAGCCCTCC	481
Qy	202	CAGGTAGCTGAG	ACTATATGTG-CTGCTACCATGCACAGCTGATTTTAAATTTTTTTG	260
Db	480	CAAGTAGCTGG	ACTACAGGTGTGCGCACCATGCCAGCTAATTTTGTATTTTGTGTA	421
Qy	261	TAGAGATGG-----	AGTTGCCAGGGTGGTCTTGAACCTCCGGCCTGAGGTGATCCT	312
Db	420	GAAACAAGGTTT	CCCCCATGTTGCCCAGGCTGCTTGAACCTCCTGGGCACAGGGGATCTG	361
Qy	313	CCTCGCTTGAC	CTCCCAAGTATCTTAGACTACAGATGCATCCACACAGCT	363
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GenCore version 5.1.6
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Perfect score: 365

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Scoring table: IDENTITY NUC

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Searched: 7331713 seqs, 327154945 residues

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Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	365	100.0	10365	8	Sequence 9, Appli
2	365	100.0	36022	13	Sequence 1708, Ap
3	147.6	40.4	301692	17	Sequence 11, Appl
4	147.6	40.4	310268	19	Sequence 195, App
5	147.2	40.3	414295	20	Sequence 6876, Ap
6	146.8	40.2	209822	21	Sequence 17560, A
7	145.8	39.9	430442	20	Sequence 128, App

c 8	145.4	39.8	437	13	US-10-027-632-46474	Sequence 46474, A
c 9	145.4	39.8	437	13	US-10-027-632-46475	Sequence 46475, A
c 10	145.4	39.8	437	17	US-10-027-632-46474	Sequence 46474, A
c 11	145.4	39.8	437	17	US-10-027-632-46475	Sequence 46475, A
c 12	145	39.7	10735	20	US-10-483-241-5	Sequence 5, Appli
c 13	144.6	39.6	339	19	US-10-674-124A-25157	Sequence 25157, A
c 14	143.8	39.4	1980090	20	US-10-719-993-6815	Sequence 6815, Ap
c 15	143.8	39.4	1980090	21	US-10-741-600-17676	Sequence 17676, A
c 16	143.6	39.3	11216	10	US-09-764-872-490	Sequence 490, App
c 17	142.6	39.1	155937	20	US-10-723-860-2208	Sequence 2208, Ap
c 18	141.8	38.8	3095	13	US-10-027-632-113781	Sequence 113781, Ap
c 19	141.8	38.8	3095	17	US-10-027-632-113781	Sequence 113781, Ap
c 20	141.6	38.8	160274	22	US-10-893-315-140	Sequence 140, App
c 21	141.6	38.8	160300	22	US-10-893-315-151	Sequence 151, App
c 22	140.8	38.6	607	13	US-10-027-632-188264	Sequence 188264, A
c 23	140.8	38.6	607	17	US-10-027-632-188264	Sequence 188264, A
c 24	140.8	38.6	1980090	20	US-10-719-993-6815	Sequence 6815, Ap
c 25	140.8	38.6	1980090	21	US-10-741-600-17676	Sequence 17676, A
c 26	140.6	38.5	827	13	US-10-027-632-157392	Sequence 157392, A
c 27	140.6	38.5	827	13	US-10-027-632-157393	Sequence 157393, A
c 28	140.6	38.5	827	17	US-10-027-632-157393	Sequence 157392, A
c 29	140.6	38.5	827	17	US-10-027-632-157393	Sequence 157393, A
c 30	140.6	38.5	51917	21	US-10-741-600-17758	Sequence 17758, A
c 31	140.4	38.5	130349	21	US-10-741-600-17619	Sequence 17619, A
c 32	140.4	38.5	138363	19	US-10-367-094-117	Sequence 117, App
c 33	140.2	38.4	150573	22	US-10-981-277-56	Sequence 56, Appl
c 34	140.2	38.4	169659	19	US-10-322-696-70	Sequence 70, Appl
c 35	140	38.4	14346	17	US-10-074-024-505	Sequence 505, App
c 36	140	38.4	34739	13	US-10-087-192-1846	Sequence 1846, Ap
c 37	140	38.4	546025	20	US-10-719-993-6862	Sequence 6862, Ap
c 38	139.8	38.3	5230	15	US-10-213-948-8	Sequence 8, Appli
c 39	139.6	38.2	26928	17	US-10-374-979-6	Sequence 6, Appli
c 40	139.6	38.2	26928	18	US-10-182-936A-6	Sequence 6, Appli
c 41	139.6	38.2	26928	19	US-10-731-739-6	Sequence 6, Appli
c 42	139.6	38.2	26928	20	US-10-477-238A-6	Sequence 6, Appli
c 43	139.6	38.2	26928	20	US-10-680-287A-6	Sequence 6, Appli
c 44	139.6	38.2	26928	21	US-10-477-173-6	Sequence 6, Appli
c 45	139.6	38.2	26928	22	US-10-834-377-6	Sequence 6, Appli

ALIGNMENTS

RESULT 1

US-08-731-499-9
; Sequence 9, Application US/08731499
; Publication No. US20030148270A1
; GENERAL INFORMATION:
; APPLICANT: GRAY, Joe W.
; APPLICANT: COLLINS, Colin
; APPLICANT: HWANG, Soo-In
; APPLICANT: GODFREY, Tony
; APPLICANT: KOWBEL, David
; APPLICANT: ROWEN, Johanna
; TITLE OF INVENTION: GENES FROM THE 20q13 AMPLICON AND THEIR
; TITLE OF INVENTION: USES
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew'
; STREET: Two Embarcadero Center, 8th Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/731,499
; FILING DATE: 16-OCT-1996
; CLASSIFICATION: 435

QY 100 AAGGTCCTTCGTCATCTAGATTGGAGTGCAGCGGTCTAATCATAGCTTACTAGT 159
Db 138314 CAGAGTCTCACTCTGTCAACCGAGTGGAGTGCAGTGTGATCTCAGCTCACTGCAAC 138373
QY 160 CTTGAATTCCTGAGTTCAAGAGATCCCTTCCTGCTCAGCTTCCAGGAGTGCAGACTATA 219
Db 138374 CTCACCTCTGGGTTCAAGCAATTCCTGCTTAGCTCCCAAGTAGCTGGACTATA 138433
QY 220 TGTGTC-TGCTACCATGCACAGCTGATTTTAAATTTTGTGAGAGATGG-----A 270
Db 138434 GGGCAGCGCCACACGCGCAGCTAAATTTTGTATTTTGTAGTAGACAGAGGTTTACCA 138493
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Db 138494 GTTGGCCAGCGGCTCTTGAACCTCTGAGCTCAAGTGCCTCCACCTCGGCTCCCA 138553
QY 331 GTATCTTAGACTACAGATGCACCTCCACACGC 362
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RESULT 7

US-10-417-375-128
; Sequence 128, Application US/10417375
; Publication No. US20040219528A1
; GENERAL INFORMATION:
; APPLICANT: David W. Morris
; APPLICANT: Marc Malandro
; TITLE OF INVENTION: Novel Therapeutic Targets in Cancer
; FILE REFERENCE: 529452001600
; CURRENT APPLICATION NUMBER: US/10/417,375
; CURRENT FILING DATE: 2003-04-15
; NUMBER OF SEQ ID NOS: 176
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 128
; LENGTH: 430442
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-417-375-128

Query Match 39.9%; Score 145.8; DB 20; Length 430442;
Best Local Similarity 67.3%; Pred. No. 1.5e-29;
Matches 239; Conservative 0; Mismatches 107; Indels 9; Gaps 2;
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Db 221592 TCACCTATTCTTTTGATGTTATTTATTTATTTATTTATTTATTTATTTATTT 221651
QY 79 GTCTTACTTTATTTTACAGAAAGGCTTCTCCGTCATCTAGATTGGAGTGCAGGGT 138
Db 221652 ATTTTATTTTATTTTGGACAGAGTCTGGCTGTGTCACCCAGGCTGGAGTGCAGTGC 221711
QY 139 GTAATCATAGCTTACTAGTCTTGAATTCCTGAGTTCAAGAGATCCTCTGCTCAGCT 198
Db 221712 ACGATCTCGGCTCGCTGCAACCTCTGCTCTGGTTCAAGTGAATCTCTGCTCAGCC 221771
QY 199 TCCAGGTAGCTGAGACTATATGTGCTGCTACCATGCACAGCTGATTTTAAATTTT 258
Db 221772 TCCAGGTAGCTGGATTACAAGCACT-CCACACGCGCCAGCTAAATTTTGTATTTTAG 221830
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QY 311 CTCTCGGTTGACCTCCCAAGTATCTTAGACTACAGATGCACCTCCACACGCTTG 365
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RESULT 8

US-10-027-632-46474/c
; Sequence 46474, Application US/10027632

Publication No. US20020198371A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 46474
; LENGTH: 437
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-46474
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Best Local Similarity 68.4%; Pred. No. 1.2e-30;
Matches 232; Conservative 1; Mismatches 97; Indels 9; Gaps 2;
QY 36 TTTAACTCTTAAACCTGTGTATTTTATCTTTGATTTTGTAGTCTTACTTTATTTTA 95
Db 380 TTTTATTATTAGACATTTAGATTTTCCAAATTTTCTTTTCTTTCTTTCTTTT 321
QY 96 GAGAAAGGCTTGTCTCCGTCATCTAGATTGGAGTGCAGGGGTGTAATCATAGCTTACTG 155
Db 320 GAGACAGGCTTGTCTCTGTCCAGGCTGAGTGAGTGGTGTGATCACAGCTCACTG 261
QY 156 TAGCTCTGAATTCCTGAGTTCAAGAGATCCTTCTGCTCAGCTTCCAGGTAGCTGAGAC 215
Db 260 CAGCCTCTACTTCCAGGCTCAAGTATCTCTGCTTAGCTCCCAAGTAGCTGGAC 201
QY 216 TATATG-TGCTGCTACCATGCACAGCTGATTTTAAATTTTGTAGAGTGGAGTTG 274
Db 200 CACAGGCATGCACACACACACCTGGCTAATTTTGTATTTTGTAGAGACAGGGTCTTG 141
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US-10-027-632-46475/c
; Sequence 46475, Application US/10027632
; Publication No. US20020198371A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676

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; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
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; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
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; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
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; LENGTH: 437
; TYPE: DNA
; ORGANISM: Human
; US-10-027-632-46475

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Best Local Similarity 68.4%; Pred. No. 1.2e-30;
Matches 232; Conservative 1; Mismatches 97; Indels 9; Gaps 2;

Qy 36 TTTAACTCTTAACCTGTGATTTTATCTTTTGAATTTTGTAGTCTTACTTTATTTTAA 95
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Qy 96 GAGAAAGGGTCTTCCGTCATCTAGATTGGAGTGCAGCGGTGTAATCATAGCTTACTG 155
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Qy 156 TAGCTTTGAATTCCTGAGTTCAAGAGATCCTTCTGCTCAGCTTCCCAGGTAGCTGAGAC 215
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Qy 216 TATATG-TGCTGTACATGCACAGCTGATTTTAAATTTTGTAGAGATGGAGTTG 274
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Qy 275 -----CCAGGCTGGTCTTGAACCTCGGCTGAGGTGATCCTCTCGTTGACCTC 326
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; Publication No. US20030204075A9
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; POLYMERIZATION: Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 46475
; LENGTH: 437
; TYPE: DNA
; ORGANISM: Human
; US-10-027-632-46475

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Best Local Similarity 68.4%; Pred. No. 1.2e-30;
Matches 232; Conservative 1; Mismatches 97; Indels 9; Gaps 2;

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Db 380 TTTTATTATTAGACATTTAGATTTTCCAAATTTTCTTTTCTTTTCTTTTCTTTT 321

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Db 320 GAGACAGGGTCTTGTCTGTCAACCCAGGCTCAAGTGATCTCTCTGCTTAGCCCTCCCAAGTAGCTGGAC 261

Qy 156 TAGCTTTGAATTCCTGAGTTCAAGAGATCCTTCTGCTCAGCTTCCCAGGTAGCTGAGAC 215
Db 260 CAGCCTCRACTTCCAGGCTCAAGTGATCTCTCTGCTTAGCCCTCCCAAGTAGCTGGAC 201

Qy 216 TATATG-TGCTGTACATGCACAGCTGATTTTAAATTTTGTAGAGATGGAGTTG 274
Db 200 CACAGGCATGCACACACACCTGGCTAAATTTTGTATTTTGTAGAGACAGGGTCTTG 141

Qy 275 -----CCAGGCTGGTCTTGAACCTCGGCTGAGGTGATCCTCTCGTTGACCTC 326
Db 140 CTCTATTACCCAGGCTGATCTTGAACCTCTGGCTCAGGTAATCTCCACCTTGCCCTC 81

Qy 327 CCRAAGTATCTTAGACTACAGATGCACCTCCACACGCTTG 365
Db 80 CGCGGCTGCTGGGATTACAGGTGTAGCCACACACCTG 42

RESULT 11
US-10-027-632-46475/c
; Sequence 46475, Application US/10027632
; Publication No. US20030204075A9
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; POLYMERIZATION: Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
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; US-10-027-632-46475

Query Match          39.8%; Score 145.4; DB 17; Length 437;
Best Local Similarity 68.4%; Pred. No. 1.2e-30;
Matches 232; Conservative 1; Mismatches 97; Indels 9; Gaps 2;
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; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
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; US-10-027-632-46474

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Best Local Similarity 68.4%; Pred. No. 1.2e-30;
Matches 232; Conservative 1; Mismatches 97; Indels 9; Gaps 2;

Qy 36 TTTAACTCTTAACCTGTGATTTTATCTTTTGAATTTTGTAGTCTTACTTTATTTTAA 95
Db 380 TTTTATTATTAGACATTTAGATTTTCCAAATTTTCTTTTCTTTTCTTTTCTTTT 321

Qy 96 GAGAAAGGGTCTTCCGTCATCTAGATTGGAGTGCAGCGGTGTAATCATAGCTTACTG 155
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Qy 275 -----CCAGGCTGGTCTTGAACCTCGGCTGAGGTGATCCTCTCGTTGACCTC 326
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RESULT 11
US-10-027-632-46475/c
; Sequence 46475, Application US/10027632
; Publication No. US20030204075A9
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; POLYMERIZATION: Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 46475
; LENGTH: 437
; TYPE: DNA
; ORGANISM: Human
; US-10-027-632-46475

Query Match          39.8%; Score 145.4; DB 17; Length 437;
Best Local Similarity 68.4%; Pred. No. 1.2e-30;
Matches 232; Conservative 1; Mismatches 97; Indels 9; Gaps 2;
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Qy	96	GAGAAAGGCTTGTCTCCGTCACTAGATTCGGAGTGCAGCGGTGTATCATAGCTTACTG	155
Db	320	GAGACAGGCTTGTCTGTCTACCCAGAGCTGGAGTGCAGTGGTGTGATCATCAGCTCACTG	261
Qy	156	TAGTCTTTGAATTCCTGAGTTCAAAGAGATCCTTCTGCTCTCAGCTTCCCAGGTAGCTGAGAC	215
Db	260	CAGCCTCRACTTCCCAGGCTCAAGTGATCCTCTGCTTAGCCTCCCAAGTAGCTGGAC	201
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Qy	275	-----CCCAGGCTGGTCTTTGAACCTCTGGCCTGAGGTGATCCTCGTTGACCTC	326
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; GENERAL INFORMATION:			
; APPLICANT: Broggini, Massimo			
; APPLICANT: D'Incalci, Maurizio			
; TITLE OF INVENTION: Oncosuppressive Gene			
; FILE REFERENCE: 2965-187			
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; Sequence 25157, Application US/10674124A
; Publication No. US20040197797A1
; GENERAL INFORMATION:
; APPLICANT: INOKO, Hidetoshi
; APPLICANT: TAMAYA, Gen
; TITLE OF INVENTION: GENE MAPPING METHOD USING MICROSATELLITE
; TITLE OF INVENTION: GENETIC POLYMORPHISM MARKERS
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; PRIOR FILING DATE: 2000-10-30
; PRIOR APPLICATION NUMBER: JP2000-112699
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; OTHER INFORMATION: Distance between 3'-terminus of neighbour
; OTHER INFORMATION: sequence listing upward to telomere on chr
; OTHER INFORMATION: 5'-terminus of this base sequence : 12007
US-10-674-124A-25157

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: August 30, 2005, 02:54:04 ; Search time 2193 Seconds
(without alignments)
6335.366 Million cell updates/sec

Title: US-08-731-499-9_COPY_10001_10365

Perfect score: 365

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Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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4: gb_est4.*

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6: gb_est6.*

7: gb_est7.*

8: gb_est8.*

9: gb_est9.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 5	140	38.4	516	5	BU076629 im50c03.Y
C 6	139.6	38.2	1000	4	BN452708 AGENCOURT
C 7	139.4	38.2	485	7	CN263690 170004241
C 8	139.4	38.2	678	5	EX481790 DKZP686K
C 9	139.4	38.2	788	7	CK000205 AGENCOURT
C 10	139.4	38.2	837	6	CD656882 AGENCOURT
C 11	139.2	38.1	422	7	H73550 ys10h07.r1
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C 18	138	37.8	482	8	AQ133619 HS 3047 A
C 19	138	37.8	677	9	AG094571 Pan trogl
C 20	137.8	37.8	561	5	EX475065 DKZP686F
C 21	137.8	37.8	606	6	CA748785 UI-H-FT1-
C 22	137.6	37.7	598	6	CD242461 AGENCOURT
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ALIGNMENTS

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VERSION AQ512276.1 GI:4742829
KEYWORDS GSS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 511)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE 99380589
PubMed 10443764
COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieterdejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Research Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
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Class: BAC ends
High quality sequence stop: 511.
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29 137 37.5 440 1 AA129746
30 136.4 37.4 538 6 CD691041
31 136 37.3 368 1 AI590042
32 135.8 37.2 884 4 BM044361
33 135.6 37.2 699 4 BG563044
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42 134.8 36.9 584 8 AQ111520
43 134.8 36.9 640 8 AQ385850
44 134.8 36.9 877 8 AQ749458
45 134.4 36.8 338 2 AW023111

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AI590042 tr75c02.x
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BG563044 602581828
AG145061 Pan trogl
AQ315924 RPCI11-96
AQ320405 RPCI11-10
BU929449 AGENCOURT
BC033208 Homo sapi
CN289800 170004243
B32496 HS-1015-B1-
BF769916 RC3-IT001
AQ111520 CIT-HSP-2
AQ385850 RPCI11-15
AQ749458 HS_5574_A
AW023111 df49d08.Y

BC Cancer Agency, Vancouver, BC, Canada
info@bcgsc.bc.ca

Steve Jones, Sarah Barber, Mabel Brown-John, Yaron Butterfield, Andy Chan, Steve S. Chand, William Chow, Alison Cloutier, Ruth Featherstone, Malachi Griffith, Obi Griffith, Ran Guin, Nancy Liao, Kim MacDonald, Amara Masson, Mike R. Mayo, Josh Moran, Ryan Morin, Teika Olson, Diana Palmquist, Anca Petrescu, Anna Liisa Prahbu, Parvaneh Saeedi, JR Santos, Angeline Schnerch, Ursula Skalska, Duane Smailus, Jeff Stott, Miranda Tsai, George Yang, Jacquie Schein, Asim Siddiqui, Rob Holt, Marco Marra.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LINL at: <http://image.lnl.gov>
Series: IRAL Plate: 43 Row: F Column: 20
This clone was selected for full length sequencing because it passed the following selection criteria: Hexamer frequency ORF analysis
This clone has the following problem: retained intron.

FEATURES

source
1. .2225
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:4581020"
/tissue type="Eye, retinoblastoma"
/clone lib="NIH MGC 16"
/lab host="DH10B-R"
/note="Vector: pOTB7"

ORIGIN

Query Match 38.8%; Score 141.6; DB 3; Length 2225;
Best Local Similarity 71.2%; Pred. No. 3.7e-19;
Matches 218; Conservative 0; Mismatches 79; Indels 9; Gaps 2;

Qy 66 TTTCGATTTGTTAGCTTACTTATTTTATGAGAAAGGGTCTGCTCCGTCATCTAGATT 125
Db 1782 TTTTAAACAGTGTGTATTTATTTATTTTGTGAGATGGGCTTGTCTCGCGCCAGGCT 1841

Qy 126 GGAGTCAGCGGTCTAATCATAGCTACTGAGTCTTGAATTCCTGAGTTCAAGAGATCC 185
Db 1842 GGAATGAGTGGTGCAATCATAGCTTATGAGCTCGAATTCCTGGGCTCAAGCAATCC 1901

Qy 186 TTCTGCCTCAGCTTCCAGGTAGCTGAGACTATATGTG-CTGCTACCAGTCACAGCTGAT 244
Db 1902 TCCACCTTACGTTCCCAAGTAGCGGACTATAGGAGAGTGCCACCTTACCCAGCTTAT 1961

Qy 245 TTTTAAATTTTGTGTAGAGATGGA-----GTTGCCAGGCTGGTCTTGAATCTCT 296
Db 1962 TTTTGTATTTTGTCAAGACAGGGAATCCCTATGTTGCCAGGCTGGTCTTGAATCTCT 2021

Qy 297 GGCTGAGGTGATCCTCTCGGTGACCTCCCAAGTATCTTAGACTACAGATGCACTCCA 356
Db 2022 GGGCTTAAGCGATCCGCTCGCTCCGCTTTTCAAGCACTGGAATTACAGATGTGAGCCA 2081

Qy 357 CCACGC 362
Db 2082 CCACAC 2087

RESULT 4

AQ213795 535 bp DNA linear GSS 18-SEP-1998
LOCUS HS_2178_B2_A10_MR CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=2178 Col=20 Row=B, genomic survey sequence.
DEFINITION
ACCESSION AQ213795
VERSION AQ213795.1 GI:3624996
KEYWORDS GSS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 535)

AUTHORS

Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L.

TITLE

Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

JOURNAL

Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

MEDLINE

99380589

PUBMED

10449764

COMMENT

Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 2178 row: B column: 20
Class: BAC ends
High quality sequence stop: 535.

FEATURES

source
1. .535
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/clone="Plate=2178 Col=20 Row=B"
/sex="male"
/clone lib="CIT Approved Human Genomic Sperm Library D"
/note="Organ: sperm; Vector: pBelobAC11; BAC Clones in E-Coli DH10B"

ORIGIN

Query Match 38.4%; Score 140.2; DB 8; Length 535;
Best Local Similarity 70.1%; Pred. No. 9.1e-19;
Matches 216; Conservative 0; Mismatches 89; Indels 3; Gaps 2;

Qy 59 TTATCTTTGATTTGTTAGTCTTACTTTATTTTATGAGAAAGGGTCTTGTCCGTCAT 118
Db 106 TTTCTATTAGTTNTGTTTGTCTTGTGTTGTGAGACAGAGTCTTGTCTGTGTGC 165

Qy 119 CTAGATTGGATGTCAGGGGTGAATCATAGCTTACTGTAGTCTTGAATTCCTGAGTTCAA 178
Db 166 CCAGGCTGGAGTGCATATGGTGTGATCTTGGCTCACTGCAGCCCTCCGCTCCAGGTTCAA 225

Qy 179 GAGATCCTTCTGCTCAGCTTCCAGGTAGCTGAGACTATATGTG-TCGTACCATGCAC 237
Db 226 GTGATTTCTGCTGCTCAGCTACCGAGCAGCTGGGACTCAGGTGCGTGCACCACTCC 285

Qy 238 AGCTGATTTTAAATTTTGTGTAGAGATGGAGTTGCCAGGCTGGTCTTTGAACCTCTG 297
Db 286 AGCTAATTTT--GTATTTTGTAGAGACCATGTTGGCCAGGCTGGTCTCGAACTCCTG 343

Qy 298 GCCTGAGGTGATCCTCTCGGTGACCTCCCAAGTATCTTAGACTACAGATGCACTCCAC 357
Db 344 ACCTCAAGTATGCTGCTGCTCAGCTCCCAAGTCTGGGATTACAGGTGTGAGCCAC 403

Qy 358 CACGCTTG 365
Db 404 CACACCTG 411

RESULT 5

BU076629/c 516 bp mRNA linear EST 27-AUG-2002
LOCUS BU076629 im50c03.y1 HR85 islet Homo sapiens cDNA clone IMAGE:6038404 5', mRNA sequence.
DEFINITION
ACCESSION BU076629
VERSION BU076629.1 GI:22517811
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 516)

AUTHORS	Melton, D., Brown, J., Kenty, G., Permutt, A., Lee, C., Kaestner, K., Lemishka, I., Seearce, M., Brestelli, J., Gradwohl, G., Clifton, S., Hillier, L., Marra, M., Pape, D., Wylie, T., Martin, J., Blistain, A., Schmitt, A., Theising, B., Ritter, E., Ronko, I., Bennett, J., Cardenas, M., Gibbons, M., McCann, R., Cole, R., Tsagareishvili, R., Williams, T., Jackson, Y. and Bowers, Y.
TITLE	Endocrine Pancreas Consortium
JOURNAL	Unpublished (2000)
COMMENT	Contact: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue Endocrine Pancreas Consortium Harvard University, Howard Hughes Medical Institute Dept of Molecular and Cellular Biology, 7 Divinity Ave, Cambridge, MA 02138 Tel: 617-495-1812 Fax: 617-495-8557 Email: dmelton@biohp.harvard.edu Library was constructed by Dr. Hiroshi Inoue DNA sequencing by: Washington University Genome Sequencing Center For information on obtaining a clone please contact: Dr. Hiroshi Inoue (hinoue@im.wustl.edu) Seq primer: -40RP from Gibco High quality sequence stop: 468. Location/Qualifiers
FEATURES	1. .516 Location/Qualifiers
source	1. .516 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="taxon:9606" /clone="IMAGE:6038404" /tissue_type="Purified pancreatic islet" /lab_host="DH10B" /lab_lib="H985 islet" /notes="Organ: Pancreas; Vector: pBluescript SK(-); Site_1: NotI; Site_2: XhoI; cDNA made by oligo-dT priming. Size-selected on agarose gel. Average insert size ~1kb. 5' XhoI site was destroyed after directional cloning. Amplified once. Contact information: Hiroshi Inoue, MD, Metabolism Div. (Alan Permutt Lab), Washington University School of Medicine, Box 8127, 660 South Euclid Ave., St. Louis, MO 63110, E-mail: hinoue@imgate.wustl.edu, Tel: 314-362-1916, Fax: 314-747-2692."
ORIGIN	
Query Match	38.4%; Score 140; DB 5; Length 516;
Best Local Similarity	70.9%; Pred. No. 1e-18;
Matches	217; Conservative 0; Mismatches 80; Indels 9; Gaps 2;
Qy	66 TTTGATTTGTTAGTCTTACTTTATTTTATTTAGAAAGGGCTTGTCCTCGTCACTAGATT 125
Db	421 TTTTAAACAGTGTGTATTTATTTATTT

SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 678)
 AUTHORS Bahr, A., Lauber, J., Mewes, H.W., Weil, B., Amid, C., Osanger, A.,
 Fobo, G., Han, M. and Wiemann, S.
 TITLE EST (Bahr, A., Lauber, J., Mewes, H.W., Weil, B., et al.)
 JOURNAL Unpublished (2003)
 COMMENT Contact: MIPS
 MIPS
 Ingolstaedter Landstr.1, D-85764 Neuherberg, Germany
 This is the 5' sequence of the clone insert
 Clone from S. Wiemann, Molecular Genome Analysis, German Cancer
 Research Center (DKFZ); Email: s.wiemann@dkfz-heidelberg.de;
 sequenced by Qiagen (Hilden/Germany) within the cDNA sequencing
 consortium of the German Genome Project.
 No sl sequence available.
 This clone (DKFZp686K11227) is available at the RZPD in Berlin.
 Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059
 Berlin- Charlottenburg, GERMANY; Email: clone@rzpd.de.
 Location/Qualifiers
 1..678
 .organism="Homo sapiens"
 .mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="DKFZp686K11227"
 /dev_stage="adult"
 /lab_host="DH108"
 /clone_lib="686 (synonym: hlcc3)"
 /notes="Vector: pTriplex2; Site_1: SfiIA; Site_2: SfiIB;
 cDNA-collection"
 ORIGIN
 Query Match 38.2%; Score 139.4; DB 5; Length 678;
 Best Local Similarity 69.0%; Pred. No. 1.3e-18;
 Matches 223; Conservative 0; Mismatches 91; Indels 9; Gaps 2;
 QY 52 GTGTAATTTATCTTTGATTTCTTTAGTCTTACTTTATTTTATAGAAAGGGTCTTGCT 111
 Db 101 GGGTTTTTTTTGTTTTGTTTGTGTTTGTGTTTTTGTGACGGTGTCTCGCT 160
 QY 112 CGGTCACTAGATGAGTGACGGGTGTAATCATAGCTTACTGTAGTCTTGAATTCCTG 171
 Db 161 GCATCAACCCAGGCTGGAGTGCAGTGGCGCAATCTCGCTTCTCGAGTCTCCGCCCTCTG 220
 QY 172 AGTTCAAGAGATCCTTCTGCCTCAGCTTCCAGGTAGCTGAGACTATATGCTGTCTACC 231
 Db 221 AGTTCAAGCGAATTCCTCGCTCAGCTTCCCAAGTAGTGGAGTGCAGCGCGCACACCC 280
 QY 232 ATGCACAGCTGATTTTAAATTTTTTTTGTAGA-----GATGGAGTTGCCCAGGCTG 283
 Db 281 ACGCCAGCTAATTTTGTATTTTATAGTACGCGGGTTTCATCATGTTGACCGAGCTG 340
 QY 284 GTCTTGAACTCTCGGCTGAGGTGATCCTCTCGGTGA-CCTCCCAAGTATCTTAGACT 342
 Db 341 GTCTCAAACTCTCTGACTTCAGGTGATCCACCCGCCCTTCAGCCTCCCAACGTTGGGATT 400
 QY 343 ACAGATGCACCTCCACCACGCTTG 365
 Db 401 ACAGCCGTGAGCCGCGCGCCTG 423
 RESULT 9
 CK000205 LOCUS 788 bp mRNA linear EST 26-NOV-2003
 DEFINITION AGENCOURT_16363613 NIH_MGC_220 Homo sapiens cDNA clone
 IMAGE:30707474 5', mRNA sequence.
 ACCESSION CK000205
 VERSION CK000205.1 GI:38526239
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

ORIGIN

Query Match 38.2%; Score 139.4; DB 6; Length 837;
 Best Local Similarity 69.0%; Pred. No. 1.2e-18;
 Matches 223; Conservative 0; Mismatches 91; Indels 9; Gaps 2;

Qy 52 GTGATATTTTATCTTTTGGATTTGTTAGTCTTACTTATTTTATTTAGAGAAAGGCTTGGCT 111
 |||||
 Db 215 GGGTTTTTTTTTGGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTGCT 274
 |||||

Qy 112 CCCTCATCTAGATGGAGTGCAGCGGTGAATCATAGCTTACTCTAGTCTTGAATTCCTG 171
 |||||
 Db 275 GCATCACCCAGCTGGAGTGCAGGGCGCAATCTCGCTTCTTCCAGTCTCCGCTCCTG 334
 |||||

Qy 172 AGTTCAAGAGATCTTCTGCTCAGCTTCCAGGTAGCTGAGACTATATGTGTGTAC 231
 |||||
 Db 335 AGTTCAGCGATTCTCTGCTCAGCTCCCAAGTAGCTGGACTGCAGGCGGCACACC 394
 |||||

Qy 232 ATGCACAGCTGATTTTAAATTTTTTTTGTAGA-----GATGGAGTTGCCAGGCTG 283
 |||||
 Db 395 ACGCCAGCTAAATTTTGTATTTTGTATTTTGTAGAGAGCGGGTTTCATCATGTTGACCGGCTG 454
 |||||

Qy 284 GTCTTGAACCTCGGCTGAGTGATCTCTCGTTGA-CCTCCCAAGTATCTTAGACT 342
 |||||
 Db 455 GTCTCAAACTCCTGACTTCAGGTGATCCACCGGCTTTCAGCGCTCCCAACGCTGCGGANT 514
 |||||

Qy 343 ACAGATGCACTCCACCAACGCTTG 365
 |||||
 Db 515 ACAGCGTGAGCGCGCGGCTG 537
 |||||

RESULT 11

H73550 422 bp mRNA linear EST 31-OCT-1995
 LOCUS ysl0h07.r1 Soares fetal liver spleen INFLS Homo sapiens cDNA clone
 DEFINITION IMAGE:214429 5' similar to contains Alu repetitive element; mRNA
 sequence.

ACCESSION H73550.1 GI:1046609

VERSION H73550

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;

AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 422)

Hallier.L., Lennon.G., Becker.M., Bonaldo.M.F., Chiapelli.B.,

Chisoe.S., Dietrich.N., DuBuque.T., Favello.A., Gish.W.,

Hawkins.M., Hultman.M., Kucaba.T., Lacy.M., Le.M., Le.N.,

Mardis.E., Moore.B., Morris.M., Parsons.J., Prange.C., Rifkin.L.,

Rohlfing.T., Schellenberg.K., Soares.M.B., Tan.F., Thierry-Mieg.J.,

Trevaakis.E., Underwood.K., Wohldmann.P., Waterston.R., Wilson.R.

and Marra.M.

Generation and analysis of 280,000 human expressed sequence tags

Genome Res. 6 (9), 807-828 (1996)

97044478

8889549

COMMENT Contact: Wilson RK

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

Insert Size: 487

High quality sequence stops: 368

Source: IMAGE Consortium, LBNL

This clone is available royalty-free through LLNL; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

Insert Length: 487 Std Error: 0.00

Seq primer: M13p1

High quality sequence stop: 368.

Location/Qualifiers

1. .422

/organism="Homo sapiens"

FEATURES

source

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:214429"

/sex="male"

/dev_stage="20 week-post conception fetus"

/lab_host="DH10B (ampicillin resistant)"

/notes="Organ: Liver and Spleen; Vector: pT73D (Pharmacia)

with a modified polylinker; Site 1: Pac I; Site 2: Eco RI;

1st strand cDNA was primed with a Pac I - oligo(dT) primer

[5' AACTGGAAGATTAATTAAGATCTTTTTTTTTTTTTTTT 3'],

double-stranded cDNA was ligated to Eco RI adaptors

(Pharmacia), digested with Pac I and cloned into the Pac I

and Eco RI sites of the modified pT73 vector. Library

went through one round of normalization. Library

constructed by Bento Soares and M.Fatima Bonaldo."

ORIGIN

Query Match 38.1%; Score 139.2; DB 7; Length 422;
 Best Local Similarity 69.5%; Pred. No. 1.5e-18;
 Matches 221; Conservative 0; Mismatches 88; Indels 9; Gaps 2;

Qy 57 TTTTATCTTTTGTATTTTGTAGTCTTACTTTTATTTTAGAGAAAGGCTCTGCTCGTC 116
 |||||
 Db 63 TGTCAATAATTTTTTTTCTTTCTTTCTTTTCTTTTGTAGACAGAGCTTGTCTGTGTC 122
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Qy 117 ATCTAGATGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTTGAATTCCTGAGTTC 176
 |||||
 Db 123 ACTCAGCTGGAGTGCAGTGGTGGATCTTGGCTCACTACAGCTCTGCCCTCTGAGTTC 182
 |||||

Qy 177 AAGAGATCCTTTCGCTCAGCTTCCAGGTAGCTGAGACTATATGTGCTGCTACCATGTC 235
 |||||
 Db 183 AAGCTATTCTCATGTCTCAGCTCTCTGAGTAGTGGAGCTACAGGTGCACGCCACACGC 242
 |||||

Qy 236 ACAGCTGATTTTAAATTTTTTTGTAGAGATGG-----AGTTGCCAGAGCTGTCT 287
 |||||
 Db 243 CTGGCTAAATTTTGTATTTTGTAGAGATGGGGTTTCACTAGTTGGCCAGGCTGTCT 302
 |||||

Qy 288 TGAATCTCTGGCTGAGGTGATCCTCTGCTGAGCTCCCAAGTATCTTAGACTACAGA 347
 |||||
 Db 303 CAAATCTCTGACCTCAAGTATCCACTGCTTGGCTCCCAAGTCTGGGATTACAGG 362
 |||||

Qy 348 TGCATCTCCACCCAGCTTG 365
 |||||
 Db 363 TGTGAGCCACCCAGCGCTG 380
 |||||

RESULT 12

AW884394/c

LOCUS

DEFINITION

AW884394

ACCESSION

KEYWORDS

SOURCE

ORGANISM

Homo sapiens

Homo sapiens

Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 384)

Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,

Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,

Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H.,

Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V.,

O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and

Simpson,A.J.

Shotgun sequencing of the human transcriptome with ORF expressed

sequence tags

Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

20202663

10737800

COMMENT

Contact: Simpson A.J.G.

Laboratory of Cancer Genetics

Ludwig Institute for Cancer Research

Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL
(<http://www.ludwig.org.br/scripts/gethtml2.pl?tbl=at&tbl=QV3-OT0065-290>)
300-137-a06&tc3=2000-03-29&tt4=1)
Seq primer: puc 18 forward
High quality sequence start: 10
High quality sequence stop: 384.

```

FEATURES
source
Location/Qualifiers
1..384
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_lib="OT0065"
/note="Organ: ovary; Vec:
Small; A mini-library was
from ORESTES PCR (0.5:
196,716 - Ludwig instit
into the pUC 18 vector.
mRNA and cDNA amplificat
stringency conditions."
```

	ORIGIN	strangency conditions.
Query Match	38.0%;	Score 138.8; DB 2; Length 384;
Best Local Similarity	72.6%;	Pred. No. 1.9e-18;
Matches 212;	Conservative 0;	Mismatches 67; Indels 13; Gaps 2;
Qy	87	TTATTTTTAGAAAAGGGTCTTGTCCTCGGTCAATCTAGATTGAGATGCACGGGTGAATCAT 146
Db	331	TTCTGTTTTAGACAGAGGTCTCGCTCTGTTGCTCAGGCCGAAGTGCAGTGGTGCAAATCAT 272
Qy	147	AGCTTAGCTAGTACTTCAATTCCTGAGTCCAAGAGATCCCTTCTGCGCTCAGCTCCCAGGT 206
Db	271	AGCTCACTGACAGCCTCCAACTCTCTGGGCTCAAGCGATCCTCTGCGCTCAGCCTCCCCAGT 212
Qy	207	AGCTGAGACTATATGTGCTGC-----TACCATGCACAGCTGATTTTAAAAATTTTTTGTGA 262
Db	211	AGCTGAGACTACATAGGGTGCATGACCCCATGCCCTGGCTAATTTTTHAAATTTTTTGTAAA 152
Qy	263	GAGATGG-----AGTTGCCCAGGCTGGTCTTTGAACTCTGGCCTGAGGTGAATCCTC 313
Db	151	GACAGGGGTCTCACTATGTTGCCCAGGCTGTCTCAAACCTCTTGGCCTCATGTGATCCTC 92
Qy	314	CTGGGTTGACCTCCCAAGTACTTTAGACTACAGATGCACCTCCACACGCTTG 365
Db	91	TTGCGTTGGTCTGCGCAAGTAGCTGGTACTATAGGTGTGCACCCACCACTAG 40

RESULT 13	
HSMB06754/c	
LOCUS	2198 bp mRNA linear HTC 03-AUG-2004
DEFINITION	Homo sapiens mRNA; CDNA DKFZP686A24102 (from clone DKFZP686A24102).
ACCESSION	BX641053
VERSION	BX641053.1 GI:34364972
KEYWORDS	HTC.
SOURCE	Homo sapiens (human)
ORGANISM	Homo sapiens
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE	1 (Bases 1 to 2198)
AUTHORS	Poustka,A., Albert,R., Moosmayer,P., Schupp,I., Wellenreuther,R., Mewes,H.W., Well,B., Amid,C., Osanger,A., Fobo,G., Han,M. and Wiemann,S.
CONSTRM	The German cDNA Consortium
TITLE	Direct Submission
JOURNAL	Submitted (13-JUL-2004) MIPS, Ingolstaedter Landstr.1, D-85764 Neuerberg, GERMANY
COMMENT	Cla from S. Wiemann, Molecular Genome Analysis, German Cancer

Research Center (DKFZ); Email: s.wiemann@dkfz-heidelberg.de; sequenced by DKFZ (German Cancer Research Center, Heidelberg/Germany) within the CDNA sequencing consortium of the German Genome Project. This clone (DKFZp686A24102) is available at the RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH in Berlin, Germany. Please contact RZPD for ordering: <http://www.rzpd.de/cgi-bin/products/cl.cgi?cloneID=DKFZp686A24102>. Further information about the clone and the sequencing project is available at <http://mips.gsf.de/projects/cdna/>.

```

FEATURES
source
1. 2198
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mrna"
/db_xref="taxon:9606"
/clone="DKFZp686A24102"
/tissue_type="fetal kidney"
/clone_lib="686"
synonym: h
DH10B; sites SfIIa + SfIIb
/dev stage="fetal"
/note="unspliced mrna"

```

ORIGIN

Query Match	38.0%;	Score 138.8;	DB 3;	Length 2198;
Best Local Similarity	69.0%;	Pred. No. 1.4e-18;		
Matches 225; Conservative	0;	Mismatches 87;	Indels 14;	Gaps 2;
Qy	51	TG TGTATTTTATTCTTTTGATTTGGTTTAGTCCTACTTTATTTTTTGTAGAGAAGGGCTCTTGC	110	
Dd	1246	TGACTGTCCTCATTTTATTTTATAATTTATTTATTTTATTTTGTAGACGGGGCTCTAC	1187	
Qy	111	TCCGTCATCTAGATTTGGAGTGCAGCGGTGAATCATAGCTTACTGTAGTCTTGAATTCCT	170	
Dd	1186	TCTGTCAACCAGCGCTGGAGTGCAGTGGTGGATCGTAGCTCAGTCGAGCCTCAAACCTCT	1127	
Qy	171	GAGTTCAAGAGATCTTCTGCTCAGCTTCCCAGGTAGCTGAGACTATATGTGC - TGCTA	229	
Dd	1126	GGGCTCAAGTAATCCTCTGCCTCAGCCTCCCCAAGTAGCTGGGACTATAGGTACATGTCA	1067	
Qy	230	CCATGCACAGCTGATTTTAAATTTTTTTTGTAG-----AGATGGAGTTGCC	276	
Dd	1066	CCATGCCCGCTCAATTA AAAAATTTTTTTTTTGTATAGATCGGGCTCTCACTGTATTTGAC	1007	
Qy	277	CAGGCTGGTCTTGAACTCTCTGGCGTGAAGGTGATCCTCTCGGTTTGACCTCCCAAGTATCT	336	
Dd	1006	CAGGCTGGTCTTAAACTCTGGTCTCAAGAGATCCTCTCGCTGGGCTCTCTAATATGCC	947	
Qy	337	TAGACTACAGATGCACCTCCACCAAGC	362	
Dd	946	GGCATTACAGCATGAGCCACCAAGC	921	

RESULT 14

CR749384	8056 bp	linear	HTC 19-AUG-2004
LOCUS	8056 bp	linear	HTC 19-AUG-2004
DEFINITION	Homo sapiens mRNA; cDNA DKFZP686F02110 (from clone DKFZP686F02110) .		
ACCESSION	CR749384		
VERSION	CR749384.1	GI:51476491	
KEYWORDS	HTC.		
SOURCE	Homo sapiens (human)		
ORGANISM	Homo sapiens		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
AUTHORS	1 (bases 1 to 8056) Koehler, K., Beyer, A., Mewes, H.W., Weil, B., Amid, C., Osanger, A., Fobo, G., Han, M. and Wiemann, S.		
CONSRSTM	The German cDNA Consortium		
TITLE	Direct Submission		
JOURNAL	Submitted (17-AUG-2004) MIPS, Ingolstaedter Landstr.1, D-85764 Neuherberg, GERMANY		
COMMENT	Clone from S. Wiemann, Molecular Genome Analysis, German Cancer Research Center (DKFZ); Email s.wiemann@dkfz-heidelberg.de; sequenced by BMFZ (Biomedical Research Center at the Heinrich-Heine-University, Duesseldorf/Germany) within the cDNA		

High quality sequence vector: 233:

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